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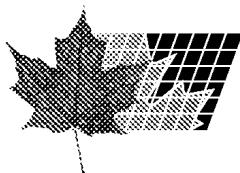
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(54) ANTICORPS RECOMBINES ANTI-GPIIB/IIIA
(54) ANTI-GPIIB/IIIA RECOMBINANT ANTIBODIES

(57)

The invention relates to novel nucleic acid sequences which code for human auto-antibodies and anti-idiotypic antibodies against blood platelet membrane proteins. The invention also relates to new amino acid sequences of human antibodies and to the use thereof in the diagnosis and therapy of diseases.



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(54) **ANTICORPS RECOMBINES ANTI-GPIIB/III A**

(54) **ANTI-GPIIB/III A RECOMBINANT ANTIBODIES**

(57) L'invention concerne de nouvelles séquences d'acide nucléique qui codent pour des auto-anticorps et des anticorps anti-idiotypes humains contre la protéine membranaire d'agrégation plaquettaire, de nouvelles séquences aminoacides d'anticorps humains et leur utilisation pour le diagnostic et la thérapie de maladies.

(57) The invention relates to novel nucleic acid sequences which code for human auto-antibodies and anti-idiotypic antibodies against blood platelet membrane proteins. The invention also relates to new amino acid sequences of human antibodies and to the use thereof in the diagnosis and therapy of diseases.



Abstract

The invention relates to novel nucleic acid sequences which encode human autoantibodies and antiidiotypic antibodies against blood platelet membrane proteins, to novel amino acid sequences of human antibodies, and to their use for the diagnosis and therapy of diseases.

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RECOMBINANT ANTI-GPIIB/IIIA ANTIBODIES

DESCRIPTION

5 The invention relates to novel nucleic acid sequences which encode human autoantibodies against blood platelet membrane proteins and which encode antiidiotypic antibodies, to novel amino acid sequences of human antibodies, and to their use for the diagnosis
10 and therapy of diseases.

Autoimmune thrombocytopenic purpura (AITP) is an immune disease which is defined by a low blood platelet count associated with normal or elevated megakaryocytopoiesis. The destruction of platelets in the reticuloendothelial system (spleen, liver and bone marrow) is increased due to the presence of anti-platelet autoantibodies. These autoantibodies, which can be detected in about 75% of AITP patients, are predominantly directed against the platelet membrane glycoproteins (GP) IIb/IIIa and Ib/IX. Several different autoantibody specificities may be found in one and the same patient (cf., e.g., Berchtold and Wenger, Blood 81 (1993), 1246-1250; Kiefel et al., Br. J. Haematol. 79 (1991), 256-262; McMillan et al., Blood 70 (1987), 1040 and Fujisawa et al., Blood 79 (1991); 1441). However, it is still difficult to characterize binding epitopes and to ascertain the pathogenic significance of the autoantibodies due to the limited 20 quantity of autoantibodies which can be obtained from AITP patients. It has only been possible to obtain a few human monoclonal antibodies from lymphocytes of AITP patients which react with GPIIb/IIIa AITP using the hybridoma technique (Kunicki et al., Hum. Antibodies Hybridomas 30 1 (1990) 83-95).

Natural autoantibodies against various selfantigens, for example against intracellular and cytoskeletal

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components of human platelets, have also been reported to occur in healthy individuals (Guilbert et al., J. Immunol. 128 (1982), 2779-2787; Hurez et al., Eur. J. Immunol. 23 (1993), 783-789 and Pfueller et al., Clin. 5 Exp. Immunol. 79 (1990), 367-373). Some of these autoantibodies which have been observed in sera from healthy individuals can also be directed against platelet-membrane proteins (Souberbielle, Eur. J. Haematol. 56 (1996), 178-180). However, the role of 10 these natural autoantibodies, and there relationship to disease-associated autoantibodies, is still unknown.

Corticosteroids can be used for treating AITP. About half of the patients react within 4 weeks to an 15 administration of prednisone; however long-term remissions are only rarely seen. The administration of high doses of intravenous immunoglobulin (IVIgG) is recommended as an emergency treatment for patients who are exhibiting severe bleeding or extremely low 20 platelet counts. This treatment is followed in most patients by a rapid, but usually only transient, increase in the platelet count. The mechanisms by which corticosteroids and IVIgG act in the treatment of AITP are still unknown. Investigations carried out by 25 Berchtold et al., (Blood 74 (1989), 2414-2417 and Berchtold and Wenger, Blood 81 (1993), 1246-1250) have disclosed that antiidiotypic antibodies which are present in IVIgG can inhibit the binding of autoantibodies to platelet glycoproteins.

30 The problem underlying the present application is that of identifying novel DNA sequences which are responsible for autoantibodies binding to GPIIb/IIIa. This approach can be used for making available novel 35 pharmaceutical preparations which can be employed for improving the diagnosis and therapy of AITP.

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It was surprisingly possible to identify binding sequences from autoantibodies after using peripheral circulating B cells from a healthy human donor to prepare a combinatorial phagemid display library of 5 human antibody heavy and light chains. Following the presentation of human heavy and light antibody Fab fragments on the surface of the filamentous phage M13, it was possible to identify phage clones which exhibit specific binding to GPIIb/IIIa.

10

For this, the phagemid library was brought consecutively into contact with thrombasthenic platelets lacking GPIIb/IIIa (negative selection) and normal platelets (positive selection). After several 15 rounds of selection and amplification by infecting *E.coli*, 23 clones were obtained which were able to bind to the GPIIb/IIIa complex. Inhibition studies using pools of monoclonal antibodies directed against the GPIIb/IIIa yielded two groups of clones: both groups 20 were inhibited by monoclonal antibodies which were specific for the GPIIb/IIIa complex and one group was also inhibited by a GPIIb-specific monoclonal antibody. These findings were confirmed by carrying out a DNA analysis of the clones which indicated the presence of 25 2 different anti-GPIIb/IIIa phage clones. These results demonstrate that 2 GPIIb/IIIa-specific phage clones, i.e. autoantibodies, can be cloned from the genome of a healthy individual and that these clones are able to recognize confirmational epitopes belonging to the 30 GPIIb/IIIa complex. Inhibition studies furthermore established that both phage clones inhibit the binding of platelet-associated autoantibodies from AITP patients to purified GPIIb/IIIa and therefore presumably recognize GPIIb/IIIa epitopes which are 35 AITP-associated. Since the phage clones contain the antigen-binding sequences of natural autoantibodies which are derived from the genome of a healthy individual, this finding can lead to new insights into

the origin of platelet-associated autoantibodies in AITP.

In addition to this, it is possible to use the novel
5 phage clones to produce recombinant antiidiotypic antibodies against anti-GPIIb/IIIa autoantibodies, with the anti-GPIIb/IIIa phage clones being used as antigen. The recombinant antiidiotypic antibodies which can be obtained in this way constitute an attractive clinical
10 alternative to using IVIgG.

The nucleotide sequences of the identified phage clones, and the amino acid sequences which are deduced from these nucleotide sequences, are depicted in the
15 sequencing listings SEQ ID No. 1 to 8 (autoantibodies) and SEQ ID No. 9 to 18 (antiidiotypic antibodies).

I. Autoantibodies

20 A first aspect of the present invention relates to nucleic acids which encode auto-antibodies. Part of the subject-matter of the invention is therefore a nucleic acid which encodes the heavy chain of a human antibody, or a functional derivative or a fragment thereof, and
25 encompasses a CDR3 region, selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

V L P F D P I S M D V, (I)

(b) a nucleotide sequence which encodes the amino acid sequence:

A L G S W G G W D H Y M D V, (II)

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a) or (b), and

(d) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to GPIIb/IIIa.

The novel nucleic acid furthermore preferably comprises a CDR1 region selected from:

5 (a) a nucleotide sequence which encodes the amino acid sequence:

G Y S W R, (III)

(b) a nucleotide sequence which encodes the amino acid sequence:

10 S Y A M H, (IV)

and

15 (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a) or (b).

15

The novel nucleic acid preferably furthermore comprises a CDR2 region selected from:

20 (a) a nucleotide sequence which encodes the amino acid sequence:

D I S Y S G S T K Y K P S L R S, (V)

(b) a nucleotide sequence which encodes the amino acid sequence:

25 V I S Y D G S N K Y Y A D S V K G, (VI)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably of at least 90%, with an amino acid sequence from (a) or (b).

30 A second aspect of the present invention is a nucleic acid which encodes the light chain of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:

35 (a) a nucleotide sequence which encodes the amino acid sequence:

A T W D D G L N G P V, (VII)

(b) a nucleotide sequence which encodes the amino acid sequence:

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A A W D D S L N G W V, (VIII)

5 (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably of at least 90%, with an amino acid sequence from (a) or (b), and

(d) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to GPIIb/IIIa.

10 The novel nucleic acid preferably furthermore comprises a CDR1 region selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

S G S S S N I R S N P V S, (IX)

15 (b) a nucleotide sequence which encodes the amino acid sequence:

S G S S S N I G S N T V N, (X)

and

20 (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a) or (b).

In addition, the novel nucleic acid preferably further 25 comprises a CDR2 region selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

G S H Q R P S, (XI)

30 (b) a nucleotide sequence which encodes the amino acid sequence:

S N N Q R P S, (XII)

and

35 (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a) or (b).

II. Antiidiotypic antibodies

A second aspect of the present invention relates to nucleic acids which encode antiidiotypic antibodies.

5 Part of the subject-matter of the invention is therefore a nucleic acid which encodes the heavy chain of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:

10 (a) a nucleotide sequence which encodes the amino acid sequence:
V R D L G Y R V L S T F T F D I, (XIII)

(b) a nucleotide sequence which encodes the amino acid sequence:
15 D G R S G S Y A R F D G M D V, (XIV)

(c) a nucleotide sequence which encodes the amino acid sequence:
M G S S V V A T Y N A F D I, (XV)

(d) a nucleotide sequence which encodes the amino acid sequence:
20 D A D G D G F S P Y Y F P Y, (XVI)

(e) a nucleotide sequence which encodes the amino acid sequence:
L R N D G W N D G F D I, (XVII)

25 (f) a nucleotide sequence which encodes the amino acid sequence:
D S E T A I A A A G R F D I, (XVIII)

(g) a nucleotide sequence which encodes the amino acid sequence:
30 E D G T T V P S Q P L E F, (XIX)

(h) a nucleotide sequence which encodes the amino acid sequence:
G S G S Y L G Y Y F D Y, (XX)

(i) a nucleotide sequence which encodes the amino acid sequence:
35 G L R S Y N Y G R N L D Y, (XXI)

(j) a nucleotide sequence which encodes an amino acid sequence having an homology of at least

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80%, and preferably of at least 90%, with an amino acid sequence from (a), (b), (c) or (d), and

5 (k) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to autoantibodies against GPIIb/IIIa.

10 The novel nucleic acid furthermore preferably comprises a CDR1 region selected from: a nucleotide sequence which encodes the amino acid sequences N F A M S, S Y T M H, D Y A L H or S H Y W S shown in Tab. 7a, a nucleotide sequence which encodes the amino acid sequence T Y Y W S, a nucleotide sequence which encodes the amino acid sequences D Y G M H, S H T I S, 15 K Y A I H or E L S M H shown in Tab. 7b, and a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with one of the previously mentioned amino acid sequences.

20 Preferably, the novel nucleic acid furthermore comprises a CDR2 region selected from a nucleotide sequence which encodes the amino acid sequences G I S G G G L L T H Y A (D/N) S V K G, L I S Y D G S N K Y Y A 25 D S V K G, G I S W D S T S I G Y A D S V K G or F I Y D G A R T R F N P S L R S shown in Tab. 7a, a nucleotide sequence which encodes the amino acid sequence YIYYSGNTNYNPSLKS, a nucleotide sequence which encodes the amino acid sequences A I S Y D G S N K Y Y A D S V 30 K G, G I T P I F G T V N Y A Q K F Q G, A I S S N G G N T Y Y A D S V K G or G F D P E D G E T I Y A Q K F Q G shown in Tab. 7b, and a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably of at least 90%, with one of 35 the previously mentioned amino acid sequences.

Another part of the subject-matter of the present invention is a nucleic acid which encodes the light

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chain of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:

5 (a) a nucleotide sequence which encodes the amino acid sequence:
C S Y V H S S T N, (XXII)

10 (b) a nucleotide sequence which encodes the amino acid sequence:
Q V W D N T N D Q, (XXIII)

15 (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a), and

15 (d) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to autoantibodies against GPIIb/IIIa.

20 Preferably, the novel nucleic acid furthermore comprises a CDR1 region selected from a nucleotide sequence which encodes the amino acid sequence T G T S S A I G N Y N F V P shown in Tab. 7a, a nucleotide sequence which encodes the amino acid sequence G G Y K I G S K S V H shown in Tab. 7b, and a nucleotide sequence which encodes an amino acid sequence having an 25 homology of at least 80%, and preferably of at least 90%, with the previously mentioned amino acid sequence.

30 In addition, the novel nucleic acid preferably furthermore comprises a CDR2 region selected from a nucleotide sequence which encodes the amino acid sequence E G S K R P S shown in Tab. 7a, a nucleotide sequence which encodes the amino acid sequence E D S Y R P S shown in Tab. 7b, and a nucleotide sequence which 35 encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with the previously mentioned amino acid sequence.

- 10 -

Within the meaning of the present invention, the phrase "functional derivative of a chain of a human antibody" is to be understood as meaning a polypeptide which encompasses at least a CDR3 region of the heavy and/or light chain, as defined above, and which is able, where appropriate together with the relevant complementary chain of the human antibody (or a derivative of such a chain), to form an antibody derivative which possesses a recognition specificity for an antigen which is equivalent to that possessed by the non-derivatized antibody. Preferably, such an antibody derivative has a binding constant for the relevant antigen of at least 10^{-6} 1/mol, preferably of at least 10^{-3} 1/mol.

15 Functional derivatives of chains of a human antibody can be prepared, for example, by using recombinant DNA techniques to delete, substitute and/or insert segments of the gene encoding the relevant polypeptide.

20 Single-chain antibodies, which can, for example, be composed of the variable domains of the H and L chains or one or two H chain domains and, where appropriate a constant domain, are particularly preferred functional derivatives of antibody chains or antibodies. The 25 preparation of such constructs is described in Hoogenboom et al., Immunol. Rev. 130 (1992), 41-68; Barbas III, Methods: Companion Methods Enzymol. 2 (1991), 119 and Plückthun, Immunochemistry (1994), Marcel Dekker Inc., Chapter 9, 210-235.

30 Within the meaning of the present invention, the phrase "equivalent ability to bind" is to be understood as being a binding affinity and/or specificity, i.e. epitope recognition, which is the same as that in the 35 specifically disclosed sequences.

Another part of the subject-matter of the present invention is a vector which contains at least one copy

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of a novel nucleic acid. This vector can be a prokaryotic vector or a eukaryotic vector. Plasmids, cosmids and bacteriophages are examples of prokaryotic vectors. Such vectors are, for example, described in 5 detail in Chapters 1 to 4 in Sambrook et al., Molecular Cloning. A Laboratory Manual, 2nd edition (1989), Cold Spring Harbor Laboratory Press. A prokaryotic vector is preferably a plasmid or a phage.

10 On the other hand, the vector can also be a eukaryotic vector, e.g. a yeast vector, an insect vector (baculovirus) or a mammalian vector (plasmid vector or viral vector). Examples of eukaryotic vectors are described in Sambrook at al., loc. cit., Chapter 16, 15 and Winnacker, Gene und Klon, Eine Einführung für die Gentechnologie [Genes and clones, an introduction to genetic engineering] (1985), VCH Verlagsgesellschaft, in particular Chapters 5, 8 and 10.

20 Yet another part of the subject-matter of the present invention is a cell which expresses a novel nucleic acid, or a cell which is transformed with a novel nucleic acid or with a novel vector. The cell can be a prokaryotic cell (e.g. a Gram-negative bacterial cell, 25 in particular E.coli) or a eukaryotic cell (e.g. a yeast, plant or mammalian cell). Examples of suitable cells and methods for introducing the novel nucleic acid into such cells can be found in the above literature references.

30 Another part of the subject-matter of the present invention is a polypeptide which is encoded by a novel nucleic acid, in particular a recombinant polypeptide. Particularly preferably, the polypeptide contains the 35 variable domain of the H chain and/or L chain of a human antibody.

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Particular preference is given to a polypeptide which exhibits antibody properties and whose subunit components are a heavy chain, or a functional derivative thereof, and a light chain, or a functional derivative thereof. The polypeptide can be composed of two separate chains or be present as a single-chain polypeptide.

Yet another part of the subject-matter of the present invention is an antibody against a novel polypeptide, which antibody is directed against a region of the polypeptide which is responsible for recognizing the antigen. This antibody can be a polyclonal antiserum, a monoclonal antibody or a fragment of a polyclonal or 15 monoclonal antibody (e.g. a Fab, $F(ab)_2$, Fab' or $F(ab')_2$ fragment). The antibody is preferably directed against the CDR3 region of the heavy and/or light antibody chain of the novel polypeptide, or a region thereof. Known methods can be used to obtain such antibodies by 20 immunizing an experimental animal with a peptide or polypeptide which contains a novel CDR3 region and isolating the resulting polyclonal antibody from the experimental animal. In addition, monoclonal antibodies can be obtained by fusing an antibody-producing B cell 25 from the experimental animal with a leukaemia cell in accordance with the method of Köhler and Milstein or a further development of this method. In addition, recombinant antibodies which are directed against the CDR3 region of the novel polypeptide can also be obtained by screening a suitable phagemid library, e.g. 30 a phagemid library from a healthy human donor, with a novel polypeptide being used as the antigen.

The invention also relates to a pharmaceutical composition which comprises a nucleic acid, a vector, a polypeptide, an antibody or a cell as previously mentioned, as active component, where appropriate together with other active components and also

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pharmaceutically customary adjuvants, additives or excipients.

5 The pharmaceutical composition can be used for preparing a diagnostic or therapeutic agent. Examples of diagnostic uses are the diagnosis of AITP or of a predisposition for AITP. Another preferred diagnostic use is that of monitoring the course of the AITP disease.

10

The use of the pharmaceutical composition as a diagnostic agent can comprise, for example, detecting a B cell subpopulation which is expressing a novel polypeptide as the antibody. This antibody can be 15 detected, for example, at the nucleic acid level, e.g. by means of a nucleic-acid-hybridization assay, together with prior amplification where appropriate. On the other hand, the antibody can also be detected as to the protein level by means of an immunoassay using 20 antigens or antibodies which react specifically with the polypeptide.

Furthermore, the novel pharmaceutical composition can also be applied in the therapeutic field, in particular 25 for the prevention or therapy of AITP. This therapeutic use can, for example, be based on stimulating the production of anti-autoantibodies. For this, the novel autoantibody polypeptide can, for example, be administered to a patient, thereby eliciting and/or 30 stimulating the formation of antiidiotypic antibodies. In this connection, this administration can be effected in accordance with customary immunization protocols (Fox et al., J. Pharmacol. Exp. Ther. 279 (1996), 1000-1008; Whittum-Hudson et al., Nat. Med. 2 (1996), 35 1116-1121; Jardieu, Curr. Opin. Immunol. 7 (1995), 779-782). On the other hand, the expression of antibody genes can be inhibited specifically by administering suitable antisense nucleic acids. The novel

antiidiotypic antibody polypeptide can be administered to a patient in order to achieve direct inhibition of the autoantibody activity.

5 Investigations carried out on the novel autoantibody polypeptides have shown that these polypeptides are surprisingly able to inhibit the binding of fibrinogen to blood platelets. The novel autoantibody polypeptides and antiidiotypic antibody polypeptides can therefore
10 be employed, where appropriate in combination, as agents for modulating blood coagulation, in particular for preventing a thrombosis, for example following cardiac infarctions or strokes, or in association with venous thromboses together with lung embolisms or
15 ischaemias, etc.

Murine monoclonal antibodies, e.g. the monoclonal antibody 7E3 (cf., e.g., US patent 5,440,020) or fragments thereof (e.g. the commercially available Fab fragment ReoPro[®]), or short synthetic peptides, have hitherto been used as fibrinogen antagonists for therapeutic purposes. However, murine monoclonal antibodies and antibody fragments suffer from the disadvantage that, as a result of their immunogenicity, 25 they give rise to undesirable side reactions when used for treating human patients, while short peptides are generally degraded very rapidly. As compared with these known agents, the novel polypeptides have the advantage that they consist of amino acid sequences of human 30 origin and therefore exhibit fewer undesirable side effects than do corresponding murine antibodies or antibody fragments, and that, because of their size, they are not subjected to such rapid degradation as are peptides.

35

The invention therefore relates to the use of a novel nucleic acid, in particular a nucleic acid which encodes an autoantibody polypeptide, of a vector which

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contains this nucleic acid, of a cell which is transformed with the nucleic acid or the vector, of a polypeptide which is encoded by the nucleic acid, or of a pharmaceutical composition which comprises one or 5 more of the said substances, for preparing an agent for affecting and in particular inhibiting the binding of fibrinogen to blood platelets. Preference is given to using the agent for modulating blood coagulation, in particular for dissolving thrombi and/or for preventing 10 the formation of thrombi. The administration of the novel pharmaceutical composition can be effected in accordance with protocols which have already been established for murine antibodies or antibody fragments.

15

Yet another part of the subject-matter of the invention is a process for isolating phagemid clones which express nucleic acids which encode autoantibodies against GPIIb/IIIa or encode antiidiotypic antibodies 20 which are directed against these autoantibodies, characterized in that a phagemid library is prepared from lymphocytes from a human donor and the desired phagemid clones are isolated by affinity selection, comprising negative and positive selection steps. 25 Preferably, the process also involves isolating antibody-encoding nucleic acids from the clones and/or using the antibody-encoding nucleic acids for expressing recombinant antibody chains or derivatives or fragments thereof.

30

The invention is also explained by the following examples, figures and sequence listings, in which

SEQ ID No. 1 shows the nucleotide sequence of the H 35 chain of a novel antibody (phagemid clone PDG7), with framework region (FR)1 extending from bp 1 to 90, complement-determining region (CDR)1 from bp 91 to

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105, FR2 from bp 106 to 147, CDR2 from bp 148 to 195, FR3 from bp 196 to 291, CDR3 from bp 292 to 324 and FR4 from bp 325 to 357,

5

SEQ ID No.2 shows the amino acid sequence corresponding to the nucleotide sequence depicted in SEQ ID No.1, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 65, FR3 from AA 66 to 97, CDR3 from AA 98 to 108 and FR4 from AA 109 to 119,

15 SEQ ID No.3 shows the nucleotide sequence of the L chain of a novel polypeptide (phagemid clone PDG7), with FR1 extending from bp 1 to 60, CDR1 from bp 61 to 99, FR2 from bp 100 to 144, CDR2 from bp 145 to 165, FR3 from bp 166 to 261, CDR3 from bp 262 to 294 and FR4 from bp 295 to 333,

20 SEQ ID No.4 shows the amino acid sequence corresponding to the nucleotide sequence given in SEQ ID No. 3, with FR1 extending from AA 1 to 20, CDR1 from AA 21 to 33, FR2 from AA 34 to 48, CDR2 from AA 49 to 55, FR3 from AA 56 to 87, CDR3 from AA 88 to 98 and FR4 from AA 99 to 11 [sic],

25 SEQ ID No.5 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone PDG13), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 109, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 294, CDR3 from

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bp 295 to 336 and FR4 from bp 337 to 369,

5 SEQ ID No.6 shows the amino sequence corresponding to the nucleotide sequence depicted in SEQ ID No.5, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CD2 from AA 50 to 66, FR3 from AA 67 to 98, CDR3 from AA 99 to 112 and FR4 from AA 113 to 123,

10

15 SEQ ID No.7 shows the nucleotide sequence of the L chain of a novel polypeptide (phagemid clone PGD13), with FR1 extending from bp 1 to 60, CDR1 from bp 61 to 99, FR2 from bp 100 to 144, CDR2 from bp 145 to 165, FR3 from bp 166 to 261, CDR3 from bp 262 to 294 and FR4 from bp 295 to 333,

20

25 SEQ ID No.8 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 7, with FR1 extending from AA 1 to 20, CDR1 from AA 21 to 33, FR2 from AA 34 to 48, CDR2 from AA 49 to 55, FR3 from AA 56 to 87, CDR3 from AA 88 to 98 and FR4 from AA 99 to 111,

30

35 SEQ ID No.9 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-X16), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 288, CDR3 from bp 289 to 336 and FR4 from bp 337 to 369,

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SEQ ID No.10 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 9, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 96, CDR3 from AA 97 to 112 and FR4 from AA 113 to 123,

5

SEQ ID No. 11 shows the nucleotide sequence of the L chain of a novel polypeptide (phagemid clone AI-X16), with FR1 extending from bp 1 to 60, CDR1 from bp 61 to 102, FR2 from bp 103 to 147, CDR2 from bp 148 to 168, FR3 from bp 169 to 264, CDR3 from [lacuna] 265 to 291 and FR4 from bp 292 to 375,

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SEQ ID No. 12 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 11, with FR1 extending from AA 1 to 20, CDR1 from AA 21 to 34, FR2 from AA 35 to 49, CDR2 from AA 50 to 56, FR3 from AA 57 to 88, CDR3 from AA 89 to 97 and FR4 from AA 89 to 125,

20

25

SEQ ID No. 13 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-X20), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 195, FR3 from bp 196 to 291, CDR3 from bp 292 to 333 and FR4 from bp 334 to 366,

30

35

SEQ ID No. 14 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 13, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA

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36 to 49, CDR2 from AA 50 to 65, FR3 from AA 66 to 97, CDR3 from AA 98 to 111 and FR4 from AA 112 to 122,

5 SEQ ID No. 15 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-X39), with FR extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from pb [sic] 148 to 198, FR3 from bp 199 to 294, CDR3 from bp 295 to 339 and FR4 from 340 to 372,

10
15 SEQ ID No. 16 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 15, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 98, CDR3 from AA 99 to 113 and FR 4 from AA 114 to 124,

20
25 SEQ ID No. 17 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-X40), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 297, CDR3 from bp 298 to 339 and FR4 from bp 340 to 372,

30
35 SEQ ID No. 18 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 17, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 99, CDR3 from AA 100 to 113 and FR4 from AA 114 to 124,

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SEQ ID No. 19 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-X2), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 195, FR3 from bp 196 to 291, CDR3 from bp 292 to 327 and FR4 from bp 328 to 360,

SEQ ID No. 20 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 19, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 65, FR3 from AA 66 to 97, CDR3 from AA 98 to 109 and FR4 from AA 110 to 120,

SEQ ID No. 21 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-B14), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 294, CDR3 from bp 295 to 336 and FR4 from bp 337 to 369;

The following variations in the sequence were also found: a C can be present at position 7, while a G can be present at position 9, a G at position 13, a G at position 15, an A at position 91, a G at position 92, a C at position 98, a T at position 149, an A at position 205, an A at position 228, an A at position 251, a T at position 253 and/or an A at position 284. The consequence of this is that, in the amino acid sequence (cf. SEQ ID No. 22), a Q can be present at

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position 3, while a V can be present at position 5, an S at position 31, an A at position 33, a V at position 50, a T at position 69, a K at position 76, an N at position 84, an S at position 85 and/or a Y at position 95.

10

SEQ ID No. 22 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 21, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 98, CDR3 from AA 99 to 112 and FR4 from AA 113 to 123,

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20

SEQ ID No. 23 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-B18), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 294, CDR3 from bp 295 to 333 and FR4 from bp 334 to 366;

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30

35

The following variations in the nucleotide sequence were also found: thus, a C can be present at position 7, while a G can be present at position 13, a C at position 16, an A at position 56, a T at position 94, a G at position 97, a T at position 155, a C at position 173, a T at position 223, a T or a C at position 252, a G at position 261, a G at position 267, an A at position 271, a C at position 275 and/or a G at position 277. The consequence of this is that, in the corresponding amino acid sequence (cf. SEQ ID No. 24), a Q can be present

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at position 3, while a V can be present at position 5, a Q at position 6, a K at position 19, a Y at position 32, an A at position 33, an I at position 52, an A at position 58, an S at position 75, an S at position 84, an R at position 87, an E at position 89, a T at position 91, an A at position 92 and/or a V at position 93.

10

SEQ ID No. 24 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 23, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 98, CDR2 from AA 99 to 111 and FR4 from AA 112 to 122,

20

25

SEQ ID No. 25 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-B24), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 294, CDR3 from bp 295 to 330 and FR4 from bp 331 to 363;

30

35

The following variations in the nucleotide sequence were also found: a C can be present at position 7, while a G can be present at position 9, a G at position 13, a G at position 15, a G at position 31, an A at position 46, a G at position 67, a G at position 89, a G at position 92, a C at position 93, a G at position 98, a G at position 102, a G at position 140, a G at position 141, a G at position 145, a T at position 149, a

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T at position 157, an A at position 158, a G at position 160, an A at position 166, an A at position 173, a T at position 235, an A at position 251, a C at position 290 and/or an A at position 293. The consequence of this is that, in the corresponding amino acid sequence (cf. SEQ ID No. 26), a Q can be present at position 3, while a V can be present at position 5, a V at position 11, an R at position 16, an A at position 23, an S at position 30, an S at position 31, a G at position 33, an M at position 34, a W at position 47, an A at position 49, a V at position 50, a Y at position 53, a D at position 54, an S at position 56, a K at position 58, an L at position 79, an N at position 84, an A at position 97 and/or a K at position 98.

20

25

SEQ ID No. 26 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 25, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 98, CDR3 from AA 99 to 110 and FR4 from AA 111 to 121,

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35

SEQ ID No. 27 shows the nucleotide sequence of the L chain of a novel polypeptide (phagemid clone AI-B24), with FR1 extending from bp 1 to 60, CDR1 from bp 61 to 96, FR2 from bp 97 to 138, CDR2 from bp 139 to 159, FR3 from bp 160 to 255, CDR3 from bp 256 to 282 and FR4 from bp 283 to 366;

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The following variations in the nucleotide sequence were also found: a C or a T can be present at position 4, while a G can be present at position 37, an A at position 40, a G at position 50, an A at position 67, a T at position 72, an A at position 133, a T at position 136, a T or a C at position 138, a G at position 148, a T at position 160, a T at position 161, a T or a C at position 162, a C at position 200, a T at position 217, a G at position 218, an A or a C at position 220, a G at position 269, a T at position 271, a G at position 272, a G at position 275 and/or a T or a C at position 282. The consequence of this is that, in the corresponding amino acid sequence (cf. SEQ ID No. 28), an I can be present at position 2, while a G can be present at position 13, a K at position 14, an R at position 17, an N at position 23, an N at position 24, an I at position 45, a Y at position 47, a D at position 50, an F at position 54, a T at position 67, an S at position 73, an R at position 74, an S at position 90, an S at position 91, an S at position 92 and/or an H at position 94.

SEQ ID No. 28 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 27, with FR1 extending from AA 1 to 20, CDR1 from AA 21 to 32, FR2 from AA 33 to 46, CDR2 from AA 47 to 53, FR3 from AA 54 to 85, CDR3 from AA 86 to 94 and FR4 from AA 95 to 122.

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SEQ ID No. 29 shows the nucleotide sequence of the H chain of a novel polypeptide (phagemid clone AI-B38), with FR1 extending from bp 1 to 90, CDR1 from bp 91 to 105, FR2 from bp 106 to 147, CDR2 from bp 148 to 198, FR3 from bp 199 to 294, CDR3 from bp 295 to 333 and FR4 from bp 334 to 366;

The following variations in the nucleotide sequence were also found: a C can be present at position 7, while a G can be present at position 9, a G at position 13, an A at position 15 and/or a C at position 16. The consequence of this is that, in the corresponding amino acid sequence, a Q can be present at position 3, while a V can be present at position 5 and/or a Q can be present at position 6, and

SEQ ID No. 30 shows the amino acid sequence of the nucleotide sequence depicted in SEQ ID No. 29, with FR1 extending from AA 1 to 30, CDR1 from AA 31 to 35, FR2 from AA 36 to 49, CDR2 from AA 50 to 66, FR3 from AA 67 to 98, CDR3 from AA 99 to 111 and FR4 from AA 112 to 122.

Figure 1 shows the inhibition of the binding of autoantibody phabs (PDG-X) to GPIIb/IIIa which is brought about by adding the antiidiotypic antibody phab AI-X17.

Figure 2 shows the inhibition of the binding of autoantibody phabs (PDG-B) to blood platelets which is brought about by antiidiotypic antibody phabs AI-B.

Figure 3 shows the binding of autoantibody phabs to untreated and EDTA-treated blood platelets,

5 Figure 4 shows the inhibition of the binding of fibrinogen to GPIIb/IIIa which is brought about by autoantibody phabs,

10 Figures 5-7 show the inhibition of the binding of autoantibody phabs to GPIIb/IIIa which is brought about by the antibody 7E3 and the antibody fragment ReoPro®.

Examples

15

1. Identification of autoantibody sequences

1.1. Isolation of autoantibodies

20 Autoantibodies were obtained from 12 AITP patients (8 suffering from primary AITP, 3 suffering from AITP associated with SLE, 1 suffering from AITP associated with Sjögren's syndrome) by incubating patient plasma with purified GPIIb/IIIa at 4°C overnight and 25 subsequently eluting, at room temperature for 15 min, in 0.2 mol/l glycine and 0.15 mol/l NaCl, pH 2.5. After centrifuging at 100,000 g for 30 min, the supernatant was neutralized with 1 mol/l Tris-HCl and dialysed overnight against Tris-buffered salt solution (TBS).

30

At the time of plasma withdrawal, all the patients were thrombocytopenic (platelet count < 150 × 10⁹/l) and had normal or enlarged megakaryocytes in the bone marrow and were free of other detectable forms of 35 immunothrombocytopenia.

1.2. Isolation of purified antigens

The antigens used were purified GPIIb/IIIa, a cytoplasmic fragment of GPIIIa (amino acids 721-744) 5 and an extracellular fragment of GPIIIa (amino acids 468-690) (Beardsley, Blut 59 (1989), 47-51 and Phillips et al., Methods Enzymol. 215 (1992), 244-263).

10 **1.3. Isolation of platelets for panning and immunoblotting**

Platelet-enriched plasma was prepared by differential centrifugation from EDTA-anticoagulated blood samples taken from healthy human donors. The platelets were 15 isolated by centrifuging at 2000 g for 15 min, then washed six times in citric acid buffer (pH 6.2) containing 50 mmol/l sodium citrate, 100 mmol/l NaCl and 125 mmol/l glucose, and finally resuspended in the same buffer.

20 The same enrichment protocol was used to obtain thrombasthenic platelets from a 14-year-old boy suffering from Glanzmann's type I thrombasthenia.

25 **1.4. Monoclonal antibodies**

Use was made of murine monoclonal antibodies which recognize the complexed form of GPIIb/IIIa and of antibodies which recognize GPIIb or GPIIIa selectively. 30 These antibodies were isolated by means of customary immunization protocols using the corresponding antigens and are not AITP-associated. The isolation of such antibodies is described in Kouns et al. (J. Biol. Chem. 267 (1992), 18844-18851), Steiner et al. (Biochim. 35 Biophys. Acta 1119 (1992), 12-21) and Häring et al. (Proc. Natl. Acad. Sci. USA 82 (1985), 4837-4841).

1.5. Phagemid library

A combinatorial Fab library was prepared in accordance with the method described by Vogel et al. (Eur. J. Immunol. 24 (1994), 1200-1207) using peripheral blood lymphocytes obtained from a healthy, preimmunized human donor. All the enzymes and oligonucleotides were obtained from Boehringer Mannheim GmbH (Mannheim, Germany) apart from the Taq polymerase (Perkin Elmer, NJ, USA). The primers for amplifying the H and L chains of the Fab molecules by PCR, the VCSM13 helper phage, and the Escherichia coli strain XL-Blue were obtained from Stratacyte (La Jolla, CA, USA). The phagemid pComb3 was obtained from Scripps Research Institute (La Jolla, CA, USA). The cloning, the transformation into XL-Blue cells and the preparation of phabs were carried out as described by Barbas III and Lerner, Methods: Companion Methods Enzymol. 2 (1991), 119). The phabs were precipitated with 4% (w/v) polyethylene glycol 8000 and 3% (w/v) NaCl and resuspended in PBS, pH 7.4. The resulting expression library contains 1×10^7 specificities.

1.6. Isolation of GPIIb/IIIa-specific phabs

GPIIb/IIIa-specific phabs were prepared by means of a total of 5 rounds of an affinity selection ("panning"). Following preabsorption (negative selection) with 5×10^7 thrombasthenic platelets, the phabs were 30 incubated for 45 min with 10^8 normal platelets (positive selection). Bound phabs were then eluted with 0.05 mol/l sodium citrate, pH 2.5, and neutralized with 1 mol/l Tris buffer. After each round of panning, the enrichment of GPIIb/IIIa-specific phabs was monitored 35 by titrating the phage-colony-forming units. After five rounds of selection, the eluted phabs were found to have been enriched by a factor of more than 100.

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The pool of phabs obtained after the fourth round of selection was analysed more closely for its GPIIb/IIIa specificity. For this, 40 phab clones were selected at random and their binding specificity was ascertained in 5 an immunodot assay. One μ l of normal and thrombasthenic platelets (10^9 ml) [sic], and also purified GPIIb/IIIa (500 μ g/ml), were added as drops onto nitrocellulose strips (Millipore Corporation, Bedford, MA, USA). The strips were blocked in TBS containing 0.15% casein 10 (TBS-casein) and then incubated overnight together with the phabs, which had been diluted in TBS-casein. After three washes with TBS-0.1% Tween 20 (TBS-Tween), the bound phabs were detected with 4-chloro-1- α -naphthol (Merck, Darmstadt, Germany) following incubation with 15 horseradish peroxidase-conjugated polyclonal rabbit anti-phage antibody (Vogel et al., loc. cit.) which had been diluted 1:1000 in TBS-casein.

20 The binding of phabs to platelets and purified GPIIb/IIIa was also tested after denaturing the proteins by heating (70°C) or by acid treatment (pH 2 with 0.5 N HCl) before dropping.

25 Of the 40 randomly selected clones, 23 (57.5%) reacted with GPIIb/IIIa, whereas 17 did not exhibit any binding. No binding of anti-GPIIb/IIIa [sic] to phabs was observed after denaturing the antigen by heat or pH 2 prior to the incubation, thereby demonstrating that intact GPIIb/IIIa is required for the phab 30 binding. Determining the presence of Fab in negative phabs revealed that 15 of the clones (88%) did not contain any Fab molecules. The two Fab-positive clones which did not bind to GPIIb/IIIa could have a low binding affinity for GPIIb/IIIa.

1.7. Fab analysis

In order to test the positive phabs for kappa (κ), lambda (λ) and Fd chains, the anti-GPIIb/IIIa phabs 5 were added as drops to nitrocellulose. The filters were incubated for 4 hours with peroxidase-labelled mouse anti-human λ , κ (The Binding Site Limited, Birmingham, England) and Fd antibodies (from the HP6045 myeloma cell line, ATCC1757, Rockville, MD, USA), which 10 antibodies had been diluted 1:1000 in TBS-casein, and then developed by chemiluminescence (ECL, Amersham, Switzerland, Zurich, Switzerland). Testing 15 randomly selected anti-GPIIb/IIIa Fab clones for κ , λ and Fd 15 chains showed that an Fd chain was present in 12 of the clones (80%) while the λ chain was present in all the clones.

Fab binding to GPIIb/IIIa on platelets was determined 20 quantitatively by preincubating pool phabs with platelets at various concentrations. The supernatant was then analysed by an immunodot method. In this connection, it was established that from 1 to 3×10^4 phabs bind per platelet. This indicates that approximately 10 to 50% of the GPIIb/IIIa molecules per 25 platelet can be occupied by phabs.

1.8. Characterizing the phab-binding epitopes

The epitope specificity of the phabs was determined by 30 carrying out an inhibition test using a variety of monoclonal antibodies (see item 4 [sic]). 1 μ l of thawed normal and thrombasthenic platelets (10^9 /ml), purified GPIIb/IIIa (500 μ g/ml), a peptide fragment of GPIIIa (amino acids 468-690, 500 μ g/ml) and the 35 cytoplasmic segment of GPIIb/IIIa (500 μ g/ml) were in each case added as drops, in duplicate, onto nitrocellulose strips. After blocking, the phab clones (0.4 μ g/ml Fab) were incubated overnight with or

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without monoclonal antibody (1 µg/ml). The bound phabs were detected using peroxidase-labelled anti-phage antibody and 4-chloro-1- α -naphthol.

5 Two groups of phab clones were identified in these investigations. While Group A (5 clones) was inhibited moderately by a pool of all the antibodies, it was inhibited strongly by GPIIb/IIIa complex-specific antibodies. Anti-GPIIb antibodies had no effect. While
10 Group B (10 clones) was inhibited completely by the pool of all the antibodies, it was inhibited to a lesser extent by the complex-specific antibody and also by the IIb-specific antibody. No group exhibited any reaction with GPIIIa-specific antibodies. The same
15 results were obtained using either platelets or purified GPIIb/IIIa as the antigen. No phab binding to the cytoplasmic peptide or to the extracellular fragment of GPIIIa was found to occur.

A summary of these results is shown in Table 1.

Table 1

Pools of monoclonal antibodies for inhibition	Inhibition of phab binding (mean value \pm SD in %)			
	Group A phab clones (n = 5)		Group B phab clones (n = 10)	
	Platelets	Purified GPIb/IIIa	Platelets	Purified GPIIb/IIIa
(1) Anti-GPIIB	0	0	49.1 \pm 5.9	49.4 \pm 9.2
(2) Anti-GPIIIa	0	0	0	0
(3) Anti-GPIIb/IIIa complex	77.8 \pm 2.9	43.6 \pm 2.1	58.6 \pm 4.4	45.5 \pm 8.0
Pool of all the antibodies	47.6 \pm 7.7	33.0 \pm 10.8	95.9 \pm 2.7	97.5 \pm 7.5
(1) - (3)				

1.9. Inhibition assays

The blocking, by the anti-GPIIb/IIIa phabs which had been found, of the binding of patient autoantibodies to 5 GPIIb/IIIa was determined by means of inhibition assays. Two of the phab clones which had been identified as previously described (PDG16 and PDG31) were used for this purpose.

10 Serial dilutions of the eluted patient autoantibodies of from 1:3 to 1:1000 were analysed for binding to purified GPIIb/IIIa. This was done by performing an immunodot assay. 100 ng of purified GPIIb/IIIa were in each case added as drops, in triplicate, onto 15 nitrocellulose strips and the filters were then blocked with TBS-casein. In order to block the binding of AITP autoantibodies to GPIIb/IIIa with phabs, the strips were incubated with 10^{11} phabs for 1 h and then incubated with varying dilutions of AITP autoantibodies 20 for 4 h. Bound autoantibodies were detected using peroxidase-labelled anti-human IgG-Fc antibodies and ECL.

25 Anti-GPIIb/IIIa phabs inhibited the binding of autoantibodies obtained from 8 AITP patients. The inhibition range [sic] was [sic] from 10 to 46%, from 32 to 60% and from 20 to 67% for PTG16, PTG31 and the pool of the two phabs, respectively. These phabs had no effect on the binding of autoantibodies obtained from 4 30 AITP patients. Both groups contained autoantibodies derived from patients suffering from primary AITP and from disease-associated AITP.

35 The results which were obtained are summarized in Table 2.

Table 2

AITP patient	Inhibition of the binding to purified GPIIB/IIIa by (%)		
	Phab clone PDG16	Phab clone PDG31	Pool of the two phab clones
WS16	13	19	40
WS37	14	20	36
KC	24	22	28
KK	22	22	40
KP	10	36	60
WS2	25	55	65
KS	60	56	64
KL	0	15	10
KG	0	0	0
KM	0	0	0
KE	0	0	0
KR	0	0	0

1.10 DNA sequence analysis

5

Plasmid DNA was purified from four Group A phab clones and 4 group [lacuna] clones using the Nukleobond® AX PC 20 purification kit (Macherey-Nagel AG, Oensingen, Switzerland).

10

The nucleic acid sequencing was carried out on an ABI373A sequencing system using a PRISM Ready Reaction DyeDeoxy Terminator Cycle Sequencing kit. The primers were obtained from Microsynth, Balgach, Switzerland.

15

The following primers were used for sequencing the H chain: Ch γ 1 (5'-CGC TGT GCC CCC AGA GGT-3') and PCH (5'-GGC CGC AAA TTC TAT TTC AAG G-3'). The following primers were used for sequencing the L chain: Cl (5'-GAG ACA CAC CAG TGT GGC-3'), Ck (5'-CAC AAC AGA GGC AGT

20

TCC-3') and PCL(5'-CTA AAC TAG CTA GTC TCC-3'). The amino acid sequences which were deduced from the DNA

sequence were compared with GenEMBL-Genbank and strain lines were assigned to VH and V λ families.

5 The VH and V λ nucleotide sequences of the 4 phab clones from each group (Group A: PDG7, PDG8, PDG10 and PDG16; Group B: PDG13, PDG17, PDG31 and PTG37 [sic]) were analysed by automated sequencing and compared with known strain line gene sequences (Tables 3 and 4). There was 100% homology in the deduced amino acid 10 sequences of the H and L chains within each group. By contrast, the homology between Group A and Group B was only 36.9% in the case of the H chain and 81.9% in the case of the L chain amino acid sequences.

15 In the H chain, Group A clones exhibit the highest degree of sequence identity with the strain line gene VH4.11 of the V_H4 family (Sanz, et al. EMBO J. 8 (1989), 3741-3748). There were 7 amino acid differences in the framework region (FR) and 8 in the complement-determining [sic] region (CDR). Group B clones differed 20 from the mostly homologous 1.9III strain line sequence of the V_H3 family (Berman et al., EMBO J. 7 (1988), 727-738) in four amino acids in the FR and one in the CDR.

25 In the L chain, the Group A and Group B clones exhibited the highest homology with the DPL2 strain line gene sequence of the V λ 1 family (Williams and Winter, Eur. J. Immunol. 323 (1993), 1456). There were 30 nine amino acid differences in FR and ten in CDR in the case of the Group A clones, and one in FR and two in CDR in the case of the Group B clones. The results which were obtained are summarized in Tables 3 and 4.

Table 3
A: Heavy chains

Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
VH4.11	QWQDQESOPGLVLUKPSSETLJLCTVSGGIS	SYH9	HIQPPGKLEHIG	YIYSGSTNNIPSLKS	RTVTSVDTSKHOFSLKLSSVNTADTVYCAR	-----	
PG1	--K-L-----W-----R-----	0-S-R	--S-----	p-S-----K-K-----R-----	-----N-----	-----	
PG10	-----	-----	-----	-----	-----	-----	
PG16	-----	-----	-----	-----	-----	-----	
1.9111	QVOLVESGGVVVQPGRLSLKSLCAGSITIS	SYGMH	WVROAPQKLENVA	VISYIGSHKYYADSVKA	RITTSRDKSKWILYLQDHSIKAEDTVYCAK	-----	
PG11	--K-L-----	--A--	-----	-----	--A-----	ALQSHGQHDLTHDV	
PG31	-----	-----	-----	-----	-----	ALQSHGQHDLTHDV	
PG33	-----	-----	-----	-----	-----	ALQSHGQHDLTHDV	
PG5155	--Q-V-----	-----	-----	-----	-----	ALQSHGQHDLTHDV	
B: Light chains							
Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
GL12	VLTQPPSASCTPGQVITSC	SASSSGENITVN	WQQLRTRKPLLY	SHHQKPS	GVPDNTFSQSKSGTGSASLATSGLQSDPENYC	ANHDDLIGQ	
PGC9	--V-----H-----	--R--P--S	--H-V-----F	GSII-----	-----R-----Q--RG-----	-T-----PV	
PGC10	-----	-----	-----	-----	-----	-----	
PGC16	-----	-----	-----	-----	-----	-----	
VL17	VLTQPPSASCTPGQVITSC	SASSSGENITVN	WQQLRTRKPLLY	SHHQKPS	GVPDNTFSQSKSGTGSASLATSGLQSDPENYC	ANHDDLIGQ	
PG11	--V-----	-----	-----	-----	-----	-----HV	
PG31	-----	-----	-----	-----	-----	-----	
PG33	-----	-----	-----	-----	-----	-----	

FR: framework region; CDR: complement-determining [sic] region. The top sequences (VH4.11; 1.9111; DPL2) are given for comparative purposes and in each case represent the deduced amino acid sequence for the most closely related published strainline gene sequence. Dashes denote identity. M85255 refers to the EMPL/GenBank reference number and denotes the deduced amino acid sequence of the human anti-GPIIb autoantibody 2E7 (Kunicki et al., J. Autoimmun. 4 (1991), 433-446). In the case of the heavy chain, the first three amino acids (QVK) are specified by the pComb3 vector sequence.

Table 4 shows the assignment of the Group A and Group B clones to known strainline V gene sequences in accordance with the amino acid homology

PDG phab clones	Heavy chain			Light chain		
	V _H family	Strain-line gene	Homology (%)	V _λ family	Strain-line gene	Homology (5)
Group A: 7, 8, 10, 16	V _H 4	V _H 4.11	84.3	V _λ I	DPL2	81.4
Group B: 13, 17, 31, 37	V _H 3	1.9III	95.1	V _λ I	DPL2	97.1

5

2. Identifying antiidiotypic antibody sequences

2.1 Phab clones AI-X

10 The phagemids technique was used to identify sequences for antiidiotypic antibodies in accordance with the method described in Example 1. The clone PDG16, which was selected in Example 1, was used as the antigen. There was no negative preselection.

15

Use was made of a pool of combinatorial phab libraries [lacuna] the specificities of a nonimmune library of peripheral B lymphocytes and of a library of peripheral lymphocytes which had been immobilized with red blood 20 cells, and also of a nonimmune library of B lymphocytes obtained from tonsils.

25 The pool of phabs which was obtained after the fourth round of panning was analysed. For this, 40 phab clones were selected at random and their binding specificities were determined. 25 of the selected clones reacted with anti-GPIIb/IIIa phab. These antiidiotypic phab clones belong to two groups: Group I (three clones) only

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reacted with Group A autoantibody phab clones (PDG 7, 8, 10 and 16), whereas the Group II phab clones (22 clones in all) reacted with the Group A and Group B phab clones, with murine monoclonal anti-GPIIb/IIIa 5 antibodies, with purified serum immunoglobulin (IVIgG) or F(ab')₂ fragments thereof, and with anti-IgE Fab. 14 phab clones (Group III) did not react with any of the substances mentioned. One Group IV phab clone only reacted with anti-GPIIb/IIIa antibodies. The results of 10 these specificity assays are summarized in Table 5a.

A DNA sequence analysis carried out on Group I phab clones (AI-X16, 17 and 24) showed complete identity in the heavy-chain-encoding sequences apart from one amino 15 acid in the CDR2 region and complete identity in the light-chain-encoding sequences. A comparison with known strainline gene sequences showed approx. 85% homology with the VH3 H chain sequence and approx. 90% homology with the V-λII L chain family sequence. A DNA sequence 20 analysis of the H chain gene was carried out on one representative of each of the Group II, III and IV phab clones. The results of this sequence analysis, and of the comparison with known strainline gene sequences, are summarized in Tables 6 and 7a.

25 The result of an inhibition assay is depicted in Fig. 1. The inhibition of the binding of AI-X17 to PDG-A by purified GPIIb/IIIa was determined by means of an immunodot assay. 660 and 220 ng of PDG-A phab, 30 respectively, were added to nitrocellulose. The antigen was incubated for 2 h with GPIIb/IIIa at concentrations in the range from 50 µg/ml to 50 ng/ml, and with a buffer solution as control, and then incubated for a further two hours with the phage clone AI-X17 (final 35 concentration 10¹²/ml). The bound phages were detected using peroxidase-conjugated polyclonal rabbit anti-phage antibody and electrochemiluminescence.

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It was found that the AI-X17 phab (Group I) is able to inhibit the binding of Group A antibody phabs (PDG-X) to the IIb/IIIa glycoprotein. This signifies that AI-X17 recognizes the antigen-binding site on PDG-A.

5

Another clone AI-X2 which binds to PDG-A was sequenced. Like clones AI-X20, 39 and 40, this clone only has a heavy chain and no light chain. The heavy chain is able to bind on its own, possibly as a dimer, to the antigen, i.e. PDG-A, with adequate specificity and affinity.

2.2 Phab clones AI-B

15 The phagemid technique was used to identify sequences of other antiidiotypic antibodies in accordance with the method described in Example 2.1. A clone PDG-B which was selected in Example 1 was used as the antigen.

20

In all, 40 phab clones were selected and their binding specificity determined. 34 of the selected clones reacted with anti-GPIIb/IIIa PHAB. These antiidiotypic phab clones belonged to three groups:

25

Group I (14 clones) only reacted with the Group B antibody phab clones, whereas the Group II phab clones (8 clones in all) reacted with both Group A and Group B phab clones. The Group III phab clones (12 clones in all) additionally reacted with murine monoclonal anti-GPIIb/IIIa antibodies, with purified serum immunoglobulin (IVIgG) or $F(ab')_2$ fragments thereof, and with anti-IgE Fab. Six phab clones (Group IV) did not react with any of the substances mentioned. The results of these specificity assays are summarized in Table 5b.

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The result of carrying out a DNA sequence analysis on Group I phab clones (AI-14, 18, 24 and 38) is summarized in Tables 6 and 7b. Clones AI-B14, 18 and 38 only had a heavy chain.

5

AI-B14 and 17 are identical. AI-B34 and 40 are likewise identical with AI-B18.

The inhibition of the binding of PDG-B to platelets by 10 AI-B phabs is depicted in Fig. 2. This was determined by means of flow-cytometric analysis. For this, a platelet-rich plasma (10^7 platelets in all) was incubated with biotinylated PDG-B in the presence or absence of AI-B phabs and using helper phages as the 15 control. The platelets were fixed with paraformaldehyde and bound PDG-B was detected with R-phycoerythrin (RPE)-labelled streptavidin. 10,000 events were counted in a FACScan appliance and the mean value of the fluorescence (\pm SD) was recorded. The strongest 20 inhibition ($> 60\%$) was achieved with clones AI-B18, 24 and 38. The inhibition of the binding shows that AI-B clones interact with the antigen-binding site on PDG-B.

Table 5a

AIX phab clones	Binding to							WO 98/55619
	PDG A	PDGB	anti-IgE Fab	anti-GPIIb/IIIa mAb	SG	F(ab') ₂		
Group I								
16,17,24	3	+	-	-	-	-	-	
Group II								
1,2,3,4,5,6,7,9,11, 13,14,23,26,27,28,29,	22	+	+	+	+	+	+	
33,35,36,37,38,40								
Group III								
8,10,12,15,18,19,21, 22,25,30,31,32,34,39	14	-	-	-	-	-	-	- 41 -
Group IV								
20	1	-	-	-	+	-	-	

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Table 5b

AI-B
phab clones

Binding to

n	phab clones	PDG-X	PDG-B	anti-IgE Fab	anti-GPIIb/IIIa mAb	IvIgG	IvIgG
14							
	(AI-B5, 7, 8, 14, 17, 18, 23, 24, 30, 31, 33, 34, 38, 40)	-	-	+	-	-	-
		-	-	-	-	-	-
8		+	+	-	-	-	-
12		+	+	+	+	+	+
6		-	-	-	-	-	-

Table 6
 anti-Id
 phage clones
 antiidiotypic
 phab clones
 (AI-X and AI-B)

	H chain			L chain		
	V _H family	Strainline gene	Homology (%) *	V _λ family	Strainline gene	Homology (%) *
AI-X16, AI-X24	V _H 3	DP47	88	V _λ 2	DPL10	88
AI-X17	V _H 3	DP47	87	V _λ 2	DPL10	88
AI-X39	V _H 3	DP49	94	-	-	-
AI-X40	V _H 3	DP31	95	-	-	-
AI-X20	V _H 4	DP71	78	-	-	-
AI-B14, AI-B17	V _H 3	DP46	91	-	-	-
AI-B18	V _H 1	DP10	85	-	-	-
AI-B24	V _H 3	DP49	81	V _λ 3	3h	82
AI-B38	V _H 1	DP5	98	-	-	-

* Highest homology (in %) of the amino acid sequences of the respective phab clones with sequences of known strainline V genes

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Table 7a

A. Heavy chains

Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
DP47	EVQLVESGGGLVQPGGSLRLSCAASGFTFS	SYNHS	HYRQAPGKGLEIVS	AISGSGSTYYADSYKQ	RITISRDHISKITYLQHISLRAEDTAVYYCAK		
AIX16	Q-K-----H-----D	HF--	-----	G---G-LI-H-----	-----H-R-V-----	VNDLGYAVLSTETFDI	MGQGTTKVYSS
AIX24	-----	-----	-----	-----	-----	-----	-----
AIX17	-----	-----	-----	-----N-----	-----	-----	-----
DP49	QVQLVESGGGLVQPGGSLRLSCAASGFTFS	SYGHH	HYRQAPGKGLEIVH	VISYDOSHKKYYADSYKQ	RITISRDHISKITYLQHISLRAEDTAVYYCAK	DGRSGSYARFDHDV	MGQGTTKVYSS
AIX39	--K-L-----H-----	--T--	-----	L-----	--A-----K-----		
DP31	EVQLVESGGGLVQPGGSLRLSCAASGFTFD	DYAHH	HYRQAPGKGLEIVS	G1SHNSGSIYADSYKQ	RITISRDHIAKISLYLQHISLRAEDTAVYYCAKD	HQSSVVATYHAFDI	MGQGTHTVYSS
AIX40	Q-K-L-----	--L-	-----	--D-T-----	-----	-----V--	
DP71	QVQLQESGPGLVKPSETLSLTCTVSGGSIS	SYWHS	HYRQAPGKGLEIIG	YIYYSSGTHHPSLKS	RVTISVDTSIQHESLKLSSVTAAEDTAVYYCAR	DADDDGFSPPYYFPY	MGQGTPVSVSS
AIX20	--K-L-----DV-R	--H--	-L-----	--DQAR-RF--R-	--SL-H-P-K-----G-----S-----		

B. Light chains

Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
DPL10	QSAITQPAVSQSGSPGQSITISC	TGTSSSDVGV9YHIVS	HYQDHPGKAPKIHIIY	EVSKRPS	GVSHRFSGSKSGHTASLTISGLQAEDEADYYC	CGYRQSSTF	
AIX16	VV-----	--A1-H--F--P-----	-----	--G-----	-----E-----	--VH---H	HVTGGGTKLTVLGQPKAAAPSVTLEPPSS
AIX24	-----	-----	-----	-----	-----	-----	-----
AIX17	-----	-----	-----	-----	-----	-----	-----

25

FR: framework region; CDR: complement-determining [sic] region. The top sequences (DP47, DP49, DP31, DP71 and DPL10) are given for comparative purposes and represent the most closely related known strainline sequences. Dashes denote identity. In the case of the heavy chain, the first three amino acids (QVK) are specified by the pComb3 vector sequence.

Table 7b

A. Heavy chains

Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
DP-16	QVQLVESGGVVQPGRSRLSCKASGFTES	SYAHII	HVRQAPGKGLEHVA	VISYDGSNHYADSVKG	RITISRDHSKNTLYLQHNSLRAEDTAVYYCAR	DSETAIAAGRFDI	HGQGTRTVTVSS
AI-B11	-K-I-.....	D-G-	-----	A-----	-----S-----H-----ST-----T-----	-----	-----
AI-B11	-----	-----	-----	-----	-----	-----
DP-10	QVQLVQSGAEVKKPGSSVKVSCKASGGTIS	SYAIS	HVRQAPGKGLEHIG	GIPPIGTANHYAQKFOG	RVTITADESTSTAYHESLSSRSEDTAVYYCAR	EDGTTVPSQLEF	HGQGTRTVTVSS
AI-B10	-K-LE-----H-----	-H-T-	-----	-----T-----V-----	-----P-----R-----T-----DDSGI-----	-----	-----
DP-49	QVQLVESGGVVQPGRSRLSCKASGFTES	SYGHII	HVRQAPGKGLEHVA	VISYDGSNHYADSVKG	RITISRDHSKNTLYLQHNSLRAEDTAVYYCAR	GSGSYLGYYIDY	HGQGTRTVTVSS
AI-B24	-K-I-----L-----Q-----S-----H-----	K-AI-	-----Y-S	A-SH-G-T-----	-----V-----S-----VR-----	-----	-----
DP-5	QVQLVQSGAEVKKPGASVKVSCKVSGYTLI	EISIII	HVRQAPGKGLEHIG	GTPDPEGETIYAQKFOG	RVTITADESTSTAYHESLSSRSEDTAVYYCAT	GIRSYHGRILD/	HGQGTRTVTVSS
AI-B30	Q-K-LE-----	-----	-----	-----	-----	-----	-----

B. Light chains

Clones	FR1	CDR1	FR2	CDR2	FR3	CDR3	FR4
VL3h	SYVLTOPIPSVSVAPGKTARIC	GGHIIIGSKSVH	H100KPGQAPVLYI	YISDRPS	FIPERESGSHSGNTATITISRVKAGREADY/C	QWIDSSSDH	
AI-B24	-V-----RQ-----T-----	-YK-----	-----V-----	C-Y-----	C-----H-----TG-----	-----H-H-Q	TIFGQGTTKLTVLRQPKAAPSVTLEPPSS

FR: framework region; CDR: complement-determining [sic] region. The top sequences (DP46, DP10, DP49, DP5 and VL3h) are given for comparative purposes and represent the most closely related known strainline sequences. Dashes denote identity. In the case of the heavy chain, the first three amino acids (QVK) are specified by the pComb3 vector sequence.

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3. Effect of autoantibody polypeptides on the binding of fibrinogen to blood platelets

3.1 Methods

5

Analysis of the binding of Fab to EDTA-pretreated blood platelets

A blood platelet-rich plasma was incubated with 10 mM 10 EDTA for 30 min. Biotinylated PDG-B and PDG-A polypeptides were added and the mixture was incubated at room temperature for 1 h. The binding of PDG-A and PDG-B to blood platelets was measured by flow-cytometric analysis using phycoerythrin-labelled 15 streptavidin.

Aggregation experiments

Blood platelet-rich plasma ($250 \times 10^9/1$) was prepared 20 freshly and maintained under 5% CO₂. The plasma was activated by different dilutions of ADP (maximum concentration 410 μ M) in the absence or presence of PDG-A or PDG-B (maximum quantity 10 μ g of Fab). The aggregation was measured in a Rodell 300BD-5 25 aggregometer (Baxter AG, D \ddot{u} dingen, Switzerland). In subsequent experiments, polyclonal anti-Fab antiserum was added to the activated platelets after PDG-A or PDG-B had been added.

30 Fibrinogen binding test

Wells of ELISA plates were coated with 0.5 μ g/ml 35 GPIIb/IIIa and blocked with 3.5% bovine serum albumin in Tris-buffered salt solution. Fibrinogen (Kabi Diagnostics, Stockholm, Sweden) was then added at different concentrations (maximally 0.08 μ g/ml) in the absence or in the presence of PDG-A, PDG-B or anti-IgE Fab as the control (maximum concentration 23 μ g/ml).

The bound fibrinogen was measured with rat anti-human fibrinogen antibody, biotinylated mouse anti-rat antibody and a conjugate consisting of streptavidin and biotinylated horseradish peroxidase (Amersham Pharmacia 5 Biotech Europe GmbH, Dübendorf, Switzerland) and using an ELISA Easy Reader (EAR340AT, SLT Instruments, Austria) at 405 nm.

10 Competition assay using the monoclonal antibody 7E3 and the antibody fragment ReoPro®

15 Platelet-rich plasma ($230 \times 10^9/l$) was incubated for 1.5 h with PDG-B or PDG-A (200 and 400 $\mu\text{g/ml}$, respectively) with or without the murine monoclonal antibody 7E3 or its Fab fragment ReoPro® (total quantity of Fab in the range from 10^{14} to 10^{10}). After fixing with an equal volume of 1% paraformaldehyde, the binding of PDG-B and PDG-A to platelets was measured by flow-cytometric analysis using phycoerythrin-labelled 20 streptavidin.

3.2 Results

25 The recombinant anti-GPIIb/IIIa Fab autoantibody fragments which were tested do not exhibit any binding to blood platelets which had been pretreated with 10 mM EDTA. This shows that the autoantibody fragments only recognize an antigen whose confirmation is intact 30 (Fig. 3).

35 In aggregation experiments using platelet-enriched plasma, neither PDG-A nor PDG-B inhibited the aggregation. In a fibrinogen-binding test in which the concentration of fibrinogen was from 10^4 to 10^6 times lower than in serum, PDG-A and PDG-B completely inhibited the fibrinogen binding (Fig. 4). No inhibition occurred when anti-IgE Fab, which had been obtained by a similar enrichment protocol, was used as

the control. These results show that both PDG-A and PDG-B interact powerfully with the fibrinogen-binding site on GPIIb/IIIa.

5 In investigations carried out with the murine monoclonal anti-GPIIb/IIIa antibody 7E3 and its commercially available Fab fragment ReoPro®, both of which inhibit the binding of fibrinogen to activated GPIIb/IIIa, the binding of PDG-B to blood platelets was
10 found to be inhibited selectively and completely (Figures 5 to 7). By contrast, the binding of PDG-A to blood platelets was not inhibited significantly either by 7E3 or by ReoPro®.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT:

5 (A) NAME:

ASAT AG Applied Science & Technology

(B) STREET: Baarerstrasse 77

(C) CITY: Zug

(E) COUNTRY: Switzerland

10 (F) POSTAL CODE: 6302

(ii) TITLE OF INVENTION: Recombinant antibodies

(iii) NUMBER OF SEQUENCES: 30

15 (iv) COMPUTER-READABLE FORM:

(A) MEDIUM TYPE: Floppy disk

(B) COMPUTER: IBM PC compatible

(C) OPERATING SYSTEM: PC-DOS/MS-DOS

20 (D) SOFTWARE: PatentIn Release #1.0,
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(vi) ORIGINAL APPLICATION DATA:

(A) APPLICATION NUMBER: DE 19723904.8

25 (B) APPLICATION DATE: 06-JUN-1997

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(A) APPLICATION NUMBER: DE 19755227.7

(B) APPLICATION DATE: 12-DEC-1997

30 (vi) ORIGINAL APPLICATION DATA:

(A) APPLICATION NUMBER: DE 19820663.1

(B) APPLICATION DATE: 08-MAY-1998

35 (2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 357 base pairs

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- (B) TYPE: nucleotide
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

5 (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) NOTATION: 1..357

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

CAG	GTG	AAA	CTG	CTC	GAG	TCG	GGC	CCA	GGA	CTG	GTG	AAG	CCT	TCG	GAG		48
Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Pro	Gly	Leu	Val	Lys	Pro	Ser	Glu		
1					5				10			15					
ACC	CTG	TCC	CTC	AAC	TGC	ACT	GTC	TCT	GGT	CGC	TCC	ATC	AGT	GGT	TAC		96
Thr	Leu	Ser	Leu	Asn	Cys	Thr	Val	Ser	Gly	Arg	Ser	Ile	Ser	Gly	Tyr		
20						25				30							
TCT	TGG	AGA	TGG	ATC	CGG	CAG	TCT	CCA	GGG	AAG	GGA	CTA	GAG	TGG	ATT		144
Ser	Trp	Arg	Trp	Ile	Arg	Gln	Ser	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Ile		
35						40				45							
GGG	GAT	ATC	TCT	TAT	AGT	GGG	AGT	ACC	AAG	TAC	AAA	CCC	TCC	CTC	AGG		192
Gly	Asp	Ile	Ser	Tyr	Ser	Gly	Ser	Thr	Lys	Tyr	Lys	Pro	Ser	Leu	Arg		
50						55				60							
AGT	CGA	GTC	ACC	CTG	TCA	GTA	GAC	ACG	TCC	AAG	AAC	CAG	TTC	TCC	CTG		240
Ser	Arg	Val	Thr	Leu	Ser	Val	Asp	Thr	Ser	Lys	Asn	Gln	Phe	Ser	Leu		
65						70				75			80				
AAG	CTG	AAT	TCG	GTG	ACC	GCT	GCG	GAC	ACG	GCC	GTC	TAT	TAC	TGT	GCG		288
Lys	Leu	Asn	Ser	Val	Thr	Ala	Ala	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
85							90					95					
CGA	GTC	TTG	CCC	TTT	GAC	CCG	ATC	TCG	ATG	GAC	GTC	TGG	GGC	AAA	GGG		336
Arg	Val	Leu	Pro	Phe	Asp	Pro	Ile	Ser	Met	Asp	Val	Trp	Gly	Lys	Gly		
100							105					110					
ACC	ACG	GTC	ACC	GTC	TCC	TCA											357
Thr	Thr	Val	Thr	Val	Ser	Ser											
			115														

(2) INFORMATION FOR SEQ ID NO: 2:

15 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 119 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

20 (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

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Gln Val Lys Leu Leu Glu Ser Gly Pro Gly Leu Val Lys Pro Ser Glu
 1 5 10 15

Thr Leu Ser Leu Asn Cys Thr Val Ser Gly Arg Ser Ile Ser Gly Tyr
 20 25 30

Ser Trp Arg Trp Ile Arg Gln Ser Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Asp Ile Ser Tyr Ser Gly Ser Thr Lys Tyr Lys Pro Ser Leu Arg
 50 55 60

Ser Arg Val Thr Leu Ser Val Asp Thr Ser Lys Asn Gln Phe Ser Leu
 65 70 75 80

Lys Leu Asn Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Val Leu Pro Phe Asp Pro Ile Ser Met Asp Val Trp Gly Lys Gly
 100 105 110

Thr Thr Val Thr Val Ser Ser
 115

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 333 base pairs

(B) TYPE: nucleotide

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..333

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

GTG GTG ACT CAG CCA CCC TCA GCG TCT GGG ACC CCC GGG CAG TGG GTC
 Val Val Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln Trp Val
 15 120 125 130 135

ACC ATC TCT TGT TCT GGG AGC AGC TCC AAC ATC AGA AGT AAT CCT GTT
 Thr Ile Ser Cys Ser Gly Ser Ser Ser Asn Ile Arg Ser Asn Pro Val
 140 145 150

AGC TGG TAT CAC CAG GTC CCA GGC ACG GCC CCC AAA CTC CTC ATC TTT
 Ser Trp Tyr His Gln Val Pro Gly Thr Ala Pro Lys Leu Leu Ile Phe
 155 160 165

GGT AGT CAT CAG CGG CCC TCA GGG GTC CCT GAC CGA TTC TCT GCC TCC
 Gly Ser His Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser Gly Ser
 170 175 180

AAG TCG GGC ACC TCC GCC TCC CTG GCC ATC CGT GGG CTC CAA TCT GGG
 Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Arg Gly Leu Gln Ser Gly
 185 190 195

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GAT GCT GGT GAC TAT TAC TGT GCA ACA TGG GAT GAC GGC CTC AAT GGT
 Asp Ala Gly Asp Tyr Tyr Cys Ala Thr Trp Asp Asp Gly Leu Asn Gly
 200 205 210 215

CCG GTG TTC GGC GGA GGG ACC AAG CTG ACC GTC CTA AGT CAG CCC
 Pro Val Phe Gly Gly Thr Lys Leu Thr Val Leu Ser Gln Pro
 220 225 230

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 111 amino acids
(B) TYPE: amino acid
(C) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

Val Val Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln Trp Val
1 5 10 15

Thr Ile Ser Cys Ser Gly Ser Ser Ser Asn Ile Arg Ser Asn Pro Val
20 25 30

Ser Trp Tyr His Gln Val Pro Gly Thr Ala Pro Lys Leu Leu Ile Phe
35 40 45

Gly Ser His Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser Gly Ser
50 55 60

Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Arg Gly Leu Gln Ser Gly
65 70 75 80

Asp Ala Gly Asp Tyr Tyr Cys Ala Thr Trp Asp Asp Gly Leu Asn Gly
85 90 95

Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Ser Gln Pro
 100 105 110

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:

15 (A) LENGTH: 369 base pairs
(B) TYPE: nucleotide
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

20 (ix) FEATURE:

(A) NAME/KEY: CDS
(B) LOCATION: 1..369

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

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CAG GTG AAA CTG CTC GAG TCT GGG GGA GGC GTG GTC CAG CCT GGG AGG	48
Gln Val Lys Leu Leu Glu Ser Gly Gly Val Val Gln Pro Gly Arg	
115 120 125	
TCC CTG AGA CTC TCC TGT GCA GCC TCT GGA TTC ACC TTC AGT AGC TAT	96
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr	
130 135 140	
GCT ATG CAC TGG GTC CGC CAG GCT CCA GGC AAG GGG CTG GAG TGG GTG	144
Ala Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val	
145 150 155	
GCA GTT ATA TCA TAT GAT GGA AGC AAT AAA TAC TAC GCA GAC TCC GTG	192
Ala Val Ile Ser Tyr Asp Gly Ser Asn Lys Tyr Tyr Ala Asp Ser Val	
160 165 170 175	
AAG GGC CGA TTC GCC ATC TCC AGA GAC AAT TCC AAG AAC ACG CTG TAT	240
Lys Gly Arg Phe Ala Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr	
180 185 190	
CTG CAA ATG AAC AGC CTG AGA GCT GAG GAC ACG GCT GTG TAT TAC TGT	288
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys	
195 200 205	
GCG AGA GCG CTG GGG AGC TGG GGG GGT TGG GAC CAC TAC ATG GAC GTC	336
Ala Arg Ala Leu Gly Ser Trp Gly Gly Trp Asp His Tyr Met Asp Val	
210 215 220	
TGG GGC AAA GGG ACC ACG GTC ACC GTC TCC TCA	369
Trp Gly Lys Gly Thr Thr Val Thr Val Ser Ser	
225 230	

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 123 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

Gln Val Lys Leu Leu Glu Ser Gly Gly Val Val Gln Pro Gly Arg	
1 5 10 15	

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr	
20 25 30	

Ala Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val	
35 40 45	

Ala Val Ile Ser Tyr Asp Gly Ser Asn Lys Tyr Tyr Ala Asp Ser Val	
50 55 60	

Lys Gly Arg Phe Ala Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr	
65 70 75 80	

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys	
85 90 95	

Ala Arg Ala Leu Gly Ser Trp Gly Gly Trp Asp His Tyr Met Asp Val	
100 105 110	

Trp Gly Lys Gly Thr Thr Val Thr Val Ser Ser	
115 120	

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(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 333 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ix) FEATURE:

10 (A) NAME/KEY: CDS
 (B) LOCATION: 1..333

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

GTG GTG ACT CAG CCA CCC TCA GCG TCT GGG ACC CCC GGG CAG AGG GTC	48
Val Val Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln Arg Val	
125 130 135	
ACC ATC TCT TGT TCT GGA AGC AGC TCC AAC ATC GGA AGT AAT ACT GTA	96
Thr Ile Ser Cys Ser Gly Ser Ser Asn Ile Gly Ser Asn Thr Val	
140 145 150 155	
AAC TGG TAC CAG CAG CTC CCA GGA ACG GCC CCC AAA CTC CTC ATC TAT	144
Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu Ile Tyr	
160 165 170	
AGT AAT AAT CAG CGG CCC TCA GGG GTC CCT GAC CGA TTC TCT GGC TCC	192
Ser Asn Asn Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser Gly Ser	
175 180 185	
AAG TCT GGC ACC TCA GCC TCC CTG GCC ATC AGT GGG CTC CAG TCT GAG	240
Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln Ser Glu	
190 195 200	
GAT GAG GCT GAT TAT TAC TGT GCA GCA TGG GAT GAC AGC CTG AAT GGT	288
Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu Asn Gly	
205 210 215	
TGG GTG TTC GGC GGA GGG ACC AAG CTG ACC GTC CTA GGT CAG CCC	333
Trp Val Phe Gly Gly Thr Lys Leu Thr Val Leu Gly Gln Pro	
220 225 230	

15 (2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 111 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

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Val Val Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln Arg Val
 1 5 10 15

Thr Ile Ser Cys Ser Gly Ser Ser Asn Ile Gly Ser Asn Thr Val
 20 25 30

Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu Ile Tyr
 35 40 45

Ser Asn Asn Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser Gly Ser
 50 55 60

Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln Ser Glu
 65 70 75 80

Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu Asn Gly
 85 90 95

Trp Val Phe Gly Gly Thr Lys Leu Thr Val Leu Gly Gln Pro
 100 105 110

(2) INFORMATION FOR SEQ ID NO: 9:

5 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 369 base pairs
- (B) TYPE: nucleotide
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

10

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..369

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

CAG GTG AAA CTG CTC GAG TCT GGG GGA GGC TTG GTT CAC CCC GGG GGG Gln Val Lys Leu Leu Glu Ser Gly Gly Gly Leu Val His Pro Gly Gly	48
115 120 125	
TCC CTG AGA CTC TCT TGT GCA GCC TCT GGA TTT ACG TTT GAC AAC TTT Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asp Asn Phe	96
130 135 140	
GCC ATG AGC TGG GTC CGC CAG GCT CCA GGG AAG GGG CTG GAG TGG GTC Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val	144
145 150 155	
TCA GGC ATT AGT GGT GGT CTT TTG ACA CAC TAC GCA GAC TCC GTG Ser Gly Ile Ser Gly Gly Leu Leu Thr His Tyr Ala Asp Ser Val	192
160 165 170 175	
AAG GGC CGG TTC ACC ATC TCC AGA AAC AAT TCC AGG AAC ACT GTA TAC Lys Gly Arg Phe Thr Ile Ser Arg Asn Asn Ser Arg Asn Thr Val Tyr	240
180 185 190	
CTA CAA ATG AAC AGC CTG AGA GCC GAA GAC ACG GCC GTG TAT TAT TGT Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys	288
195 200 205	

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GTG AGA GAT CTG GGC TAT AGA GTA CTT TCG ACT TTT ACT TTT GAT ATC 336
 Val Arg Asp Leu Gly Tyr Arg Val Leu Ser Thr Phe Thr Phe Asp Ile
 210 215 220

TGG GGC CAG GGG ACA AAG GTC ACC GTC TCT TCA 369
 Trp Gly Gln Gly Thr Lys Val Thr Val Ser Ser
 225 230

(2) INFORMATION FOR SEQ ID NO: 10:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 123 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:
 Gln Val Lys Leu Leu Glu Ser Gly Gly Gly Leu Val His Pro Gly Gly
 1 5 10 15
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asp Asn Phe
 20 25 30
 Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45
 Ser Gly Ile Ser Gly Gly Leu Leu Thr His Tyr Ala Asp Ser Val
 50 55 60
 Lys Gly Arg Phe Thr Ile Ser Arg Asn Asn Ser Arg Asn Thr Val Tyr
 65 70 75 80
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
 85 90 95
 Val Arg Asp Leu Gly Tyr Arg Val Leu Ser Thr Phe Thr Phe Asp Ile
 100 105 110
 Trp Gly Gln Gly Thr Lys Val Thr Val Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 11:

15 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 375 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

20

(ix) FEATURE:

(A) NAME/KEY: CDS
 (B) LOCATION: 1..375

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

GTG GTG ACT CAG CCT GCC TCC GTG TCT GGG TCT CCT GGA CAG TCG ATC	48
Val Val Thr Gln Pro Ala Ser Val Ser Gly Ser Pro Gly Gln Ser Ile	
125 130 135	
ACC ATC TCC TGC ACT GGA ACC AGC AGT GCT ATT GGG AAT TAT AAC TTT	96
Thr Ile Ser Cys Thr Gly Thr Ser Ser Ala Ile Gly Asn Tyr Asn Phe	
140 145 150 155	
GTC CCC TGG TAC CAA CAG CAC CCA GGC AAA GCC CCC AAA CTC ATG ATT	144
Val Pro Trp Tyr Gln Gln His Pro Gly Lys Ala Pro Lys Leu Met Ile	
160 165 170 175	
TAT GAG GGC AGT AAG CGG CCC TCA GGG GTT TCT AAT CGC TTC TCT GGC	192
Tyr Glu Gly Ser Lys Arg Pro Ser Gly Val Ser Asn Arg Phe Ser Gly	
175 180 185	
TCC AAG TCT GGC AAC ACG GCC TCC CTG ACA ATC TCT GGG CTC CAG GCT	240
Ser Lys Ser Gly Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu Gln Ala	
190 195 200	
GAG GAC GAG GCT GAG TAT TAC TGC TGC TCA TAT GTT CAT AGT AGC ACT	288
Glu Asp Glu Ala Glu Tyr Tyr Cys Cys Ser Tyr Val His Ser Ser Thr	
205 210 215	
AAT TGG GTG TTC GGC GGA GGG ACC AAG CTG ACC GTC CTA GGT CAG CCC	336
Asn Trp Val Phe Gly Gly Thr Lys Leu Thr Val Leu Gly Gln Pro	
220 225 230 235	
AAG GCT GCC CCC TCG GTC ACT CTG TTC CCA CCC TCC TCT	375
Lys Ala Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser	
240 245	

5 (2) INFORMATION FOR SEQ ID NO: 12:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 125 amino acids

(B) TYPE: amino acid

10 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

Val Val Thr Gln Pro Ala Ser Val Ser Gly Ser Pro Gly Gln Ser Ile	
1 5 10 15	
Thr Ile Ser Cys Thr Gly Thr Ser Ser Ala Ile Gly Asn Tyr Asn Phe	
20 25 30	
Val Pro Trp Tyr Gln Gln His Pro Gly Lys Ala Pro Lys Leu Met Ile	
35 40 45	
Tyr Glu Gly Ser Lys Arg Pro Ser Gly Val Ser Asn Arg Phe Ser Gly	
50 55 60	
Ser Lys Ser Gly Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu Gln Ala	
65 70 75 80	
Glu Asp Glu Ala Glu Tyr Tyr Cys Cys Ser Tyr Val His Ser Ser Thr	
85 90 95	

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Asn	Trp	Val	Phe	Gly	Gly	Gly	Thr	Lys	Leu	Thr	Val	Leu	Gly	Gln	Pro
100								105						110	
Lys	Ala	Ala	Pro	Ser	Val	Thr	Leu	Phe	Pro	Pro	Ser	Ser			
115							120					125			

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:

5

- (A) LENGTH: 366 base pairs
- (B) TYPE: nucleotide
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

10

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..366

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

CAG	GTG	AAA	CTG	CTC	GAG	TCA	GGA	CCA	GGA	CTG	GTG	AAG	CCC	TCG	GAG	48
Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Pro	Gly	Leu	Val	Lys	Pro	Ser	Glu	
130								135						140		
ACC	CTG	TCT	CTC	ACC	TGC	ACT	GTC	TCT	GAT	GTC	TCC	ATC	AGA	AGT	CAT	96
Thr	Leu	Ser	Leu	Thr	Cys	Thr	Val	Ser	Asp	Val	Ser	Ile	Arg	Ser	His	
145							150					155				
TAC	TGG	AGT	TGG	CTC	CGG	CAG	CCC	CCA	GGG	AAG	GGA	CTG	GAG	TGG	ATT	144
Tyr	Trp	Ser	Trp	Leu	Arg	Gln	Pro	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Ile	
160						165					170					
GGG	TTT	ATC	TAT	GAC	GGT	GCG	AGA	ACC	AGG	TTC	AAC	CCC	TCC	CTC	AGG	192
Gly	Phe	Ile	Tyr	Asp	Gly	Ala	Arg	Thr	Arg	Phe	Asn	Pro	Ser	Leu	Arg	
175						180				185						
AGT	CGA	GTC	TCC	CTT	TCA	ATG	GAC	CCA	TCC	AAG	AAG	CAG	TTT	TCC	CTG	240
Ser	Arg	Val	Ser	Leu	Ser	Met	Asp	Pro	Ser	Lys	Lys	Gln	Phe	Ser	Leu	
190						195			200			205				
AAA	CTG	GGG	TCT	GTG	ACC	GCT	GCG	GAC	TCG	GCC	GTC	TAC	TAC	TGT	GCG	288
Lys	Leu	Gly	Ser	Val	Thr	Ala	Ala	Asp	Ser	Ala	Val	Tyr	Tyr	Cys	Ala	
210						215						220				
AGA	GAC	GCG	GAT	GGA	GAT	GGC	TTC	AGC	CCA	TAC	TAC	T-T	CCC	TAC	TGG	336
Arg	Asp	Ala	Asp	Gly	Asp	Gly	Pro	Ser	Pro	Tyr	Tyr	Phe	Pro	Tyr	Trp	
225						230			235							
GGC	CAG	GGA	ATC	CCG	GTC	TCC	GTC	TCC	TCG							366
Gly	Gln	Gly	Ile	Pro	Val	Ser	Val	Ser	Ser							
240						245										

(2) INFORMATION FOR SEQ ID NO: 14

(i) SEQUENCE CHARACTERISTICS:

20

- (A) LENGTH: 122 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

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(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Pro	Gly	Leu	Val	Lys	Pro	Ser	Glu
1				5				10			15				

Thr	Leu	Ser	Leu	Thr	Cys	Thr	Val	Ser	Asp	Val	Ser	Ile	Arg	Ser	His
				20				25				30			

Tyr	Trp	Ser	Trp	Leu	Arg	Gln	Pro	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Ile
				35			40			45					

Gly	Phe	Ile	Tyr	Asp	Gly	Ala	Arg	Thr	Arg	Phe	Asn	Pro	Ser	Leu	Arg
				50			55			60					

Ser	Arg	Val	Ser	Leu	Ser	Met	Asp	Pro	Ser	Lys	Lys	Gln	Phe	Ser	Leu
65				70				75				80			

Lys	Leu	Gly	Ser	Val	Thr	Ala	Ala	Asp	Ser	Ala	Val	Tyr	Tyr	Cys	Ala
				85				90				95			

Arg	Asp	Ala	Asp	Gly	Asp	Gly	Phe	Ser	Pro	Tyr	Tyr	Phe	Pro	Tyr	Trp
			100				105					110			

Gly	Gln	Gly	Ile	Pro	Val	Ser	Val	Ser	Ser						
			115			120									

5 (2) INFORMATION FOR SEQ ID NO: 15:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 372 base pairs

(B) TYPE: nucleotide

10 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(ix) FEATURE:

(A) NAME/KEY: CDS

15 (B) LOCATION: 1..372

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

CAG	GTG	AAA	CTG	CTC	GAG	TCT	GGG	GGA	GGC	GTG	GTC	CAC	CCT	GGG	AGG	48
Gln	Val		Lys	Leu	Glu	Ser	Gly	Gly	Gly	Val	Val	His	Pro	Gly	Arg	
							125				130			135		

TCC	CTG	AGA	CTC	TCC	TGT	GCA	GCC	TCT	GGA	TTC	ACC	TTC	AGT	AGC	TAT	96
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	
							140			145		150				

ACT	ATG	CAC	TGG	GTC	CGC	CAG	GCT	CCA	GGC	AAG	GGG	CTG	GAG	TGG	GTG	144
Thr	Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	
							155			160		165		170		

GCA	CTT	ATA	TCA	TAT	GAT	GGG	AGC	AAT	AAA	TAC	TAC	GCA	GAC	TCC	GTG	192
Ala	Leu	Ile	Ser	Tyr	Asp	Gly	Ser	Asn	Lys	Tyr	Tyr	Ala	Asp	Ser	Val	
							175			180		185				

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AAG GGC CGA TTC GCC ATC TCC AGA GAC AAT TCC AAG AAC ACG CTA TAT	240
Lys Gly Arg Phe Ala Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr	
190	195
195	200
CTG CAA ATG AAC AGC CTG AGA GCT GAG GAC ACG GCT GTG TAT TAC TGT	288
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys	
205	210
210	215
GCG AAA GAT GGC CGG AGT GGG AGC TAC GCC AGG TTC GAC GGT ATG GAC	336
Ala Lys Asp Gly Arg Ser Gly Ser Tyr Ala Arg Phe Asp Gly Met Asp	
220	225
225	230
GTC TGG GGC CAA CGG ACC ACG GTC ACC GTC TCC TCA	372
Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser	
235	240
240	245

(2) INFORMATION FOR SEQ ID NO: 16:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 124 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

Gln Val Lys Leu Leu Glu Ser Gly Gly Val Val His Pro Gly Arg			
1	5	10	15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr			
20	25	30	
Thr Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val			
35	40	45	
Ala Leu Ile Ser Tyr Asp Gly Ser Asn Lys Tyr Tyr Ala Asp Ser Val			
50	55	60	
Lys Gly Arg Phe Ala Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr			
65	70	75	80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys			
85	90	95	
Ala Lys Asp Gly Arg Ser Gly Ser Tyr Ala Arg Phe Asp Gly Met Asp			
100	105	110	
Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser			
115	120		

(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS:

15 (A) LENGTH: 372 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

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(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..372

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

CAG	GTG	AAA	CTG	CTC	GAG	TCT	GGG	GGA	GGC	TTG	GTA	CAG	CCT	GGC	AGG	48
Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Arg	
125					130					135					140	
TCC	CTG	AGA	CTC	TCC	TGT	GCA	GCC	TCT	GGG	TTC	ACC	TTT	GAT	GAT	TAT	96
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Asp	Asp	Tyr	
					145					150					155	
GCC	CTG	CAC	TGG	GTC	CGT	CAA	GCT	CCA	GGG	AAG	GGC	CTG	GAG	TGG	GTC	144
Ala	Leu	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	
					160					165					170	
TCA	GGT	ATT	AGT	TGG	GAT	AGT	GGT	ACC	ATA	GGC	TAT	GCG	GAC	TCT	GTG	192
Ser	Gly	Ile	Ser	Trp	Asp	Ser	Gly	Thr	Ile	Gly	Tyr	Ala	Asp	Ser	Val	
					175					180					185	
AAG	GGC	CGA	TTC	ACC	ATC	TCC	AGA	GAC	AAC	GCC	AAG	AAC	TCC	CTG	TAT	240
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ala	Lys	Asn	Ser	Leu	Tyr	
					190					195					200	
CTG	CAA	ATG	AAC	AGT	CTG	AGA	GCT	GAG	GAC	ACG	GCC	TTG	TAT	TAC	TGT	288
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Leu	Tyr	Tyr	Cys	
					205					210		215			220	
GTA	AAA	GAT	ATG	GGG	TCT	TCG	GTA	GTG	GCT	ACG	TAC	AAT	GCT	TTT	GAT	336
Val	Lys	Asp	Met	Gly	Ser	Sex	Val	Val	Ala	Thr	Tyr	Asn	Ala	Phe	Asp	
					225					230					235	
ATC	TGG	GGC	CAA	GGG	ACA	ATG	GTC	ACC	GTC	TCT	TCA					372
Ile	Trp	Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser					
					240					245						

(2) INFORMATION FOR SEQ ID NO: 18:

10 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 124 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

15 (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:

Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Arg	
1					5					10					15	
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Asp	Asp	Tyr	
					20					25					30	
Ala	Leu	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	
					35					40					45	
Ser	Gly	Ile	Ser	Trp	Asp	Ser	Gly	Thr	Ile	Gly	Tyr	Ala	Asp	Ser	Val	
					50					55					60	
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ala	Lys	Asn	Ser	Leu	Tyr	
					65					70					75	

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Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Leu Tyr Tyr Cys
 85 90 95

Val Lys Asp Met Gly Ser Ser Val Val Ala Thr Tyr Asn Ala Phe Asp
 100 105 110

Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 19:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 360 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: cDNA for mRNA

(vii) IMMEDIATE SOURCE:

(B) CLONE (E): AI-X2

15 (ix) FEATURE:

(A) NAME/KEY: CDS
 (B) LOCATION: 1..360

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:

CAG GTG AAA CTG CTC GAG TCA GGC CCA GGA CTG GTG AAG CCT TCG GAG 48
 Gln Val Lys Leu Leu Glu Ser Gly Pro Gly Leu Val Lys Pro Ser Glu
 125 130 135 140

ACC CTG CTC ACC TGC ACT GTC TCT GGT GGC TCC TTC AGT ACT TAC 96
 Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Phe Ser Thr Tyr
 145 150 155

TAT TGG AGC TGG ATC CGG CAG CCC CCA GGG AAG GGA CTG GAG TGG ATT 144
 Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 160 165 170

GGG TAT ATC TAT TAC AGT GGG AAC ACC AAC TAC AAC CCC TCC CTC AAG 192
 Gly Tyr Ile Tyr Tyr Ser Gly Asn Thr Asn Tyr Asn Pro Ser Leu Lys
 175 180 185

AGT CGA GCC ACC ATA TCA GTA GAC ACG TCC AAG AAC CAG TTC TCC CTG 240
 Ser Arg Ala Thr Ile Ser Val Asp Thr Ser Lys Asn Gln Phe Ser Leu
 190 195 200

AAG CTG AGC TCT GTT ACC GCC GCA GAC ACG GCC GTA TAT TAC TGT GCG 288
 Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 205 210 215 220

AGA CTG CGT AAC GAT GGC TGG AAT GAT GGC TTT GAT ATC TGG GGC CAA 336
 Arg Leu Arg Asn Asp Gly Trp Asn Asp Gly Phe Asp Ile Trp Gly Gln
 225 230 235

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GGG ACA ATG GTC ACC GTC TCT TCA
 Gly Thr Met Val Thr Val Ser Ser
 240

360

(2) INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:

5

- (A) LENGTH: 120 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

10

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

Gln Val Lys Leu Leu Glu Ser Gly Pro Gly Leu Val Lys Pro Ser Glu
 1 5 10 15

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Phe Ser Thr Tyr
 20 25 30

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Asn Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Ala Thr Ile Ser Val Asp Thr Ser Lys Asn Gln Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Leu Arg Asn Asp Gly Trp Asn Asp Gly Phe Asp Ile Trp Gly Gln
 100 105 110

Gly Thr Met Val Thr Val Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 21

15

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 369 base pairs
- (B) TYPE: nucleotide
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: cDNA for mRNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

25

(vii) IMMEDIATE SOURCE:

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(B) CLONE (E): AI-B14

(viii) POSITION IN THE GENOME:

(A) CHROMOSOME/SEGMENT: 14

5 (B) MAP POSITION: q32.3

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..369

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:

CAG GTG AAA CTG CTC GAG TCT GGG GGA GGC GTG GTC CAG CCT GGG AGG	48
Gln Val Lys Leu Leu Glu Ser Gly Gly Val Val Gln Pro Gly Arg	
125 130 135	

TCC CTG AGA CTC TCC TGT GCA GCC TCT GGA TTC ACC TTC AGT GAC TAT	96
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asp Tyr	
140 145 150	

GGC ATG CAC TGG GTC CGC CAG GCT CCA GGC AAG GGG CTG GAG TGG GTG	144
Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val	
155 160 165	

GCA GCT ATA TCA TAT GAT GGA AGT AAC AAA TAC TAT GCA GAC TCC GTG	192
Ala Ala Ile Ser Tyr Asp Gly Ser Asn Lys Tyr Tyr Ala Asp Ser Val	
170 175 180	

AAG GGC CGA TTC TCC ATC TCC AGA GAC AAT TCC AAC AAT ACG CTA TAT	240
Lys Gly Arg Phe Ser Ile Ser Arg Asp Asn Ser Asn Asn Thr Leu Tyr	
185 190 195 200	

CTG CAA ATG AGC ACC CTG AGA GCT GAG GAC ACG GCT GTC TAT TTC TGT	288
Leu Gln Met Ser Thr Leu Arg Ala Glu Asp Thr Ala Val Tyr Phe Cys	
205 210 215	

GCG AGA GAT TCG CAA ACG GCA ATA GCG GCA GCT GGA CGG TTT GAT ATC	336
Ala Arg Asp Ser Glu Thr Ala Ile Ala Ala Gly Arg Phe Asp Ile	
220 225 230	

TGG GGC CAA GGG ACA ATG GTC ACC GTC TCT TCA	369
Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser	
235 240	

(2) INFORMATION FOR SEQ ID NO: 22:

15

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 123 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

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Gln Val Lys Leu Leu Glu Ser Gly Gly Gly Val Val Gln Pro Gly Arg
 1 5 10 15

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asp Tyr
 20 25 30

Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

Ala Ala Ile Ser Tyr Asp Gly Ser Asn Lys Tyr Tyr Ala Asp Ser Val
 50 55 60

Lys Gly Arg Phe Ser Ile Ser Arg Asp Asn Ser Asn Asn Thr Leu Tyr
 65 70 75 80

Leu Gln Met Ser Thr Leu Arg Ala Glu Asp Thr Ala Val Tyr Phe Cys
 85 90 95

Ala Arg Asp Ser Glu Thr Ala Ile Ala Ala Ala Gly Arg Phe Asp Ile
 100 105 110

Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 366 base pairs

(B) TYPE: nucleotide

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: cDNA for mRNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

15 (vii) IMMEDIATE SOURCE:

(B) CLONE(E): AI-B18

(viii) POSITION IN THE GENOME:

(A) CHROMOSOME SEGMENT: 14

20 (B) MAP POSITION: q32.3

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..366

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

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CAG GTG AAA CTG CTC GAG TCT GGG GCT GAG GTG AAG AAG CCT CGG TCC	48
Gln Val Lys Leu Leu Glu Ser Gly Ala Glu Val Lys Lys Pro Gly Ser	
125 130 135	
TCG GTG ATG GTC TCC TGC AAG GCT TCT GGA GGC ACC TTC AGC AGC CAT	96
Ser Val Met Val Ser Cys Lys Ala Ser Gly Gly Thr Phe Ser Ser His	
140 145 150 155	
ACT ATC AGC TGG GTG CCG CAG GCC CCT GGA CAA GGC CTT GAG TGG ATG	144
Thr Ile Ser Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met	
160 165 170	
GGA GGG ATC ACC CCT ATC TTT GGT ACA GTG AAC TAC GCA CAG AAG TTC	192
Gly Ile Thr Pro Ile Phe Gly Thr Val Asn Tyr Ala Gln Lys Phe	
175 180 185	
CAG GGC AGA GTC ACC ATT ACC GCG GAC GAA CCC ACG AGC ACA GCC TAC	240
Gln Gly Arg Val Thr Ile Thr Ala Asp Glu Pro Thr Ser Thr Ala Tyr	
190 195 200	
ATG GAA CTG AGG AGC CTG ACA TCT GAC GAC TCG GGC ATC TAT TAC TGT	288
Met Glu Leu Arg Ser Leu Thr Ser Asp Asp Ser Gly Ile Tyr Tyr Cys	
205 210 215	
GCG AGA GAA GAT GGC ACT ACA GTA CCA AGT CAA CCC CTT GAG TTC TGG	336
Ala Arg Glu Asp Gly Thr Thr Val Pro Ser Gln Pro Leu Glu Phe Trp	
220 225 230 235	
GGC CAG GGA ACC CGG GTC ACC GTC TCC TCT	366
Gly Gln Gly Thr Arg Val Thr Val Ser Ser	
240 245	

(2) INFORMATION FOR SEQ ID NO: 24

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 122 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

Gln Val Lys Leu Leu Glu Ser Gly Ala Glu Val Lys Lys Pro Gly Ser	
1 5 10 15	
Ser Val Met Val Ser Cys Lys Ala Ser Gly Gly Thr Phe Ser Ser His	
20 25 30	
Thr Ile Ser Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met	
35 40 45	
Gly Gly Ile Thr Pro Ile Phe Gly Thr Val Asn Tyr Ala Gln Lys Phe	
50 55 60	
Gln Gly Arg Val Thr Ile Thr Ala Asp Glu Pro Thr Ser Thr Ala Tyr	
65 70 75 80	
Met Glu Leu Arg Ser Leu Thr Ser Asp Asp Ser Gly Ile Tyr Tyr Cys	
85 90 95	
Ala Arg Glu Asp Gly Thr Thr Val Pro Ser Gln Pro Leu Glu Phe Trp	
100 105 110	
Gly Gln Gly Thr Arg Val Thr Val Ser Ser	
115 120	

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(2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 363 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA for mRNA

10

(vi) ORIGINAL SOURCE:

15 (A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

15 (B) CLONE (E): AI-B24

(viii) POSITION IN THE GENOME:

20 (A) CHROMOSOME/SEGMENT: 14
 (B) MAP POSITION: q32.3

(ix) FEATURE:

(A) NAME/KEY: CDS
 (B) LOCATION: 1..363

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

CAG	GTG	AAA	CTG	CTC	GAG	TCT	GGG	GGA	GGC	TTG	GTC	CAG	CCT	GGG	GGG	
Gln	Val	Lys	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	
125							130					135				

48

TCC	CTG	AGA	CTC	TCC	TGT	TCA	GCC	TCT	GGG	TTC	ACC	TTC	AAT	AAA	TAT	
Ser	Leu	Arg	Leu	Ser	Cys	Ser	Ala	Ser	Gly	Phe	Thr	Phe	Asn	Lys	Tyr	
140							145					150				

96

GCA	ATA	CAC	TGG	GTC	CGC	CAG	GCT	CCA	GGG	AAG	GGA	CTG	GAA	TAT	GTT	
Ala	Ile	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Tyr	Val	
155							160			165		170				

144

TCA	GCT	ATT	AGT	AGT	AAT	GGG	GGT	AAC	ACA	TAC	TAC	GCA	GAC	TCC	GTG	
Ser	Ala	Ile	Ser	Ser	Asn	Gly	Gly	Asn	Thr	Tyr	Tyr	Ala	Asp	Ser	Val	
175							180					185				

192

AAG	GGC	AGA	TTC	ACC	ATC	TCC	AGA	GAC	AAT	TCC	AAG	AAC	ACG	GTG	TAT	
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Val	Tyr	
190							195					200				

240

CTT	CAA	ATG	AGC	AGT	CTG	AGA	GCT	GAG	GAC	ACG	GCT	GTG	TAT	TAC	TGT	
Leu	Gln	Met	Ser	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	
205							210					215				

288

GTT	AGA	GGA	AGT	GGG	AGC	TAC	TTA	GGA	TAC	TAC	TTT	GAC	TAC	TGG	GGC	
Val	Arg	Gly	Ser	Gly	Ser	Tyr	Leu	Gly	Tyr	Tyr	Phe	Asp	Tyr	Trp	Gly	
220							225					230				

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CAG GGA ACC CTG GTC ACC GTC TCC TCA
 Gln Gly Thr Leu Val Thr Val Ser Ser
 235 240

363

(2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 121 base pairs
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

Gln Val Lys Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1 5 10 15

Ser Leu Arg Leu Ser Cys Ser Ala Ser Gly Phe Thr Phe Asn Lys Tyr
 20 25 30

Ala Ile His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Tyr Val
 35 40 45

Ser Ala Ile Ser Ser Asn Gly Gly Asn Thr Tyr Tyr Ala Asp Ser Val
 50 55 60

Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Val Tyr
 65 70 75 80

Leu Gln Met Ser Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
 85 90 95

Val Arg Gly Ser Gly Ser Tyr Leu Gly Tyr Tyr Phe Asp Tyr Trp Gly
 100 105 110

Gln Gly Thr Leu Val Thr Val Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:

15 (A) LENGTH: 366 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

20 (ii) MOLECULE TYPE: cDNA for mRNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

25 (vii) IMMEDIATE SOURCE:

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(B) CLONE (E): AI-B24

(viii) POSITION IN THE GENOME:

(A) CHROMOSOME/SEGMENT: 22

5 (B) MAP POSITION: q11

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..366

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

GTG GTG ACT CAG CCA CCC TCG GTG TCA GTG GCT CCA AGA CAG ACG GCC	48
Val Val Thr Gln Pro Pro Ser Val Ser Val Ala Pro Arg Gln Thr Ala	
125 130 135	
ACG ATT ACC TGT GGG GGA TAC AAG ATT GGA AGT AAA AGT GTC CAC TGG	96
Thr Ile Thr Cys Gly Gly Tyr Lys Ile Gly Ser Lys Ser Val His Trp	
140 145 150	
TAC CAA CAG AAG CCA GGC CAG GCC CCT GTA TTG GTC GTC TAT GAG GAT	144
Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Val Tyr Glu Asp	
155 160 165	
TCC TAC CGG CCC TCA GAG ATC CCT GAG CGA TTC TCT GGC TCC AAC TCT	192
Ser Tyr Arg Pro Ser Glu Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser	
170 175 180 185	
GGG AAC ATG GCC ACC CTG ACC ATC ACC GGG GTC GAA GCC GGG GAT GAG	240
Gly Asn Met Ala Thr Leu Thr Ile Thr Gly Val Glu Ala Gly Asp Glu	
190 195 200	
GCC GAC TAC TAC TGT CAG GTG TGG GAT AAT ACT AAT GAT CAG ACG ATA	288
Ala Asp Tyr Tyr Cys Gln Val Trp Asp Asn Thr Asn Asp Gln Thr Ile	
205 210 215	
TTC GGC GGA GGG ACC AAG CTG ACC GTC CTA CGT CAG CCC AAG GCT GCC	336
Phe Gly Gly Thr Lys Leu Thr Val Leu Arg Gln Pro Lys Ala Ala	
220 225 230	
CCC TCG GTC ACT CTG TTC CCG CCC TCC TCT	366
Pro Ser Val Thr Leu Phe Pro Pro Ser Ser	
235 240	

15 (2) INFORMATION FOR SEQ ID NO: 28:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 122 amino acids

(B) TYPE: amino acid

20 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

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Val Val Thr Gln Pro Pro Ser Val Ser Val Ala Pro Arg Gln Thr Ala
 1 5 10 15

Thr Ile Thr Cys Gly Gly Tyr Lys Ile Gly Ser Lys Ser Val His Trp
 20 25 30

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Val Tyr Glu Asp
 35 40 45

Ser Tyr Arg Pro Ser Glu Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser
 50 55 60

Gly Asn Met Ala Thr Leu Thr Ile Thr Gly Val Glu Ala Gly Asp Glu
 65 70 75 80

Ala Asp Tyr Tyr Cys Gln Val Trp Asp Asn Thr Asn Asp Gln Thr Ile
 85 90 95

Phe Gly Gly Thr Lys Leu Thr Val Leu Arg Gln Pro Lys Ala Ala
 100 105 110

Pro Ser Val Thr Leu Phe Pro Pro Ser Ser
 115 120

(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 366 base pairs
 (B) TYPE: nucleotide
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: cDNA for mRNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

15 (vii) IMMEDIATE SOURCE:

(B) CLONE(E): AI-B38

(viii) POSITION IN THE GENOME:

(A) CHROMOSOME SEGMENT: 14

20 (B) MAP POSITION: q32.3

(ix) FEATURE:

(A) NAME/KEY: CDS
 (B) LOCATION: 1..366

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

CAG GTG AAA CTG CTC GAG TCT GGG GCT GAG GTG AAG AAG CCT GGG GCC	48
Gln Val Lys Leu Leu Glu Ser Gly Ala Glu Val Lys Lys Pro Gly Ala	
125 130 135	
TCA GTG AAG GTC TCC TGC AAG GTT TCC GGA TAC ACC CTC ACT GAA TTA	96
Ser Val Lys Val Ser Cys Lys Val Ser Gly Tyr Thr Leu Thr Glu Leu	
140 145 150	
TCC ATG CAC TGG GTG CGA CAG GCT CCT GGA AAA GGG CTT GAG TGG ATG	144
Ser Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Met	
155 160 165 170	
GGA GGT TTT GAT CCT GAA GAT GGT GAA ACA ATC TAC GCA CAG AAA TTC	192
Gly Gly Phe Asp Pro Glu Asp Gly Glu Thr Ile Tyr Ala Gln Lys Phe	
175 180 185	
CAG GGC AGA GTC ACC ATG ACC GAG GAC ACA TCT ACA GAC ACG GCC TAC	240
Gln Gly Arg Val Thr Met Thr Glu Asp Thr Ser Thr Asp Thr Ala Tyr	
190 195 200	
ATG GAG CTG AGC AGC CTG AGA TCT GAG GAC ACG GCC GTG TAT TAC TGT	288
Met Glu Leu Ser Ser Leu Arg Ser Glu Asp Thr Ala Val Tyr Tyr Cys	
205 210 215	
GAG ACA GGT CTG AGG TCG TAC AAC TAT GGT CGT AAC CTT GAC TAT TGG	336
Glu Thr Gly Leu Arg Ser Tyr Asn Tyr Gly Arg Asn Leu Asp Tyr Trp	
220 225 230	
GGC CAG GGA ACC CTG GTC ACC GTC TCC TCA	366
Gly Gln Gly Thr Leu Val Thr Val Ser Ser	
235 240	

(2) INFORMATION FOR SEQ ID NO: 30:

5 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 122 amino acids
- (B) TYPE: amino acid
- (C) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

Gln Val Lys Leu Leu Glu Ser Gly Ala Glu Val Lys Lys Pro Gly Ala	
1 5 10 15	
Ser Val Lys Val Ser Cys Lys Val Ser Gly Tyr Thr Leu Thr Glu Leu	
20 25 30	
Ser Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Met	
35 40 45	
Gly Gly Phe Asp Pro Glu Asp Gly Glu Thr Ile Tyr Ala Gln Lys Phe	
50 55 60	
Gln Gly Arg Val Thr Met Thr Glu Asp Thr Ser Thr Asp Thr Ala Tyr	
65 70 75 80	
Met Glu Leu Ser Ser Leu Arg Ser Glu Asp Thr Ala Val Tyr Tyr Cys	
85 90 95	
Glu Thr Gly Leu Arg Ser Tyr Asn Tyr Gly Arg Asn Leu Asp Tyr Trp	
100 105 110	

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Gly Gln Gly Thr Leu Val Thr Val Ser Ser
115 120

1. Nucleic acid which encodes a heavy chain, which is able to bind to GPIb/IIa, of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:
 - (a) a nucleotide sequence which encodes the amino acid sequence:
V L P F D P I S M D V, (I)
 - (b) a nucleotide sequence which encodes the amino acid sequence
A L G S W G G W D H Y M D V, (II)
and
 - (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).
2. Nucleic acid according to Claim 1, which furthermore comprises a CDR1 region selected from:
 - (a) a nucleotide sequence which encodes the amino acid sequence:
G Y S W R, (III)
 - (b) a nucleotide sequence which encodes the amino acid sequence:
S Y A M H, (IV)
and
 - (c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).
3. Nucleic acid according to either Claim 1 or 2, which furthermore comprises a CDR2 region, selected from

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(a) a nucleotide sequence which encodes the amino acid sequence:

D I S Y S G S T K Y K P S L R S, (V)

(b) a nucleotide sequence which encodes the amino acid sequence:

V I S Y D G S N K Y Y A D S V K G, (VI)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).

4. Nucleic acid which encodes a light chain, which is able to bind to GPIIb/IIIa, of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

A T W D D G L N G P V, (VII)

(b) a nucleotide sequence which encodes the amino acid sequence

A A W D D S L N G W V, (VIII)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b),

with the proviso that when the nucleic acid encompasses a nucleotide sequence according to (b), it does not simultaneously comprise nucleotide sequences which encode the amino acid sequences SGSSSNIGSNTVN and SNNQRPS, and when the nucleic acid comprises a nucleotide sequence according to (c), it does not simultaneously comprise nucleotide sequences which encode the amino acid sequences SGSSSNIGSNTVN and RNNQRPS.

5. Nucleic acid according to Claim 4, which furthermore comprises a CDR1 region selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

S G S S S N I R S N P V S, (IX)

(b) a nucleotide sequence which encodes the amino acid sequence:

S G S S S N I G S N T V N, (X)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).

6. Nucleic acid according to Claim 4 or 5, which furthermore comprises a CDR2 region selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

G S H Q R P S, (XI)

(b) a nucleotide sequence which encodes the amino acid sequence:

S N N Q R P S, (XII)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).

7. Nucleic acid which encodes the heavy chain of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR3 region, selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

V R D L G Y R V L S T F T F D I, (XIII)

(b) a nucleotide sequence which encodes the amino acid sequence:

D G R S G S Y A R F D G M D V, (XIV)

(c) a nucleotide sequence which encodes the amino acid sequence:

M G S S V V A T Y N A F D I, (XV)

(d) a nucleotide sequence which encodes the amino acid sequence:

D A D G D G F S P Y Y F P Y, (XVI)

(e) a nucleotide sequence which encodes the amino acid sequence:

L R N D G W N D G F D I, (XVII)

(f) a nucleotide sequence which encodes the amino acid sequence:

D S E T A I A A A G R F D I, (XVIII)

(g) a nucleotide sequence which encodes the amino acid sequence:

E D G T T V P S Q P L E F, (XIX)

(h) a nucleotide sequence which encodes the amino acid sequence:

G S G S Y L G Y Y F D Y, (XX)

(i) a nucleotide sequence which encodes the amino acid sequence:

G L R S Y N Y G R N L D Y, (XXI)

(j) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% and preferably of at least 90%, with an amino acid sequence from (a), (b), (c) or (d), and

(k) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to autoantibodies against GPIIb/IIIa.

8. Nucleic acid according to Claim 7, which furthermore comprises a CDR1 and/or CDR2 region selected from a nucleotide sequence which encodes the amino acid sequences shown in Tab. 7a or b or an amino acid sequence which is at least 80% homologous thereto.

9. Nucleic acid which encodes the light chain of a human antibody, or a functional derivative or a fragment thereof, and comprises a CDR 3 region, selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

C S Y V H S S T N, (XXII)

(b) a nucleotide sequence which encodes the amino acid sequence:

Q V W D N T N D Q, (XXIII)

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80%, and preferably at least 90%, with an amino acid sequence from (a), and

(d) a nucleotide sequence which encodes an amino acid sequence having an equivalent ability to bind to autoantibodies against GPIIb/IIIa.

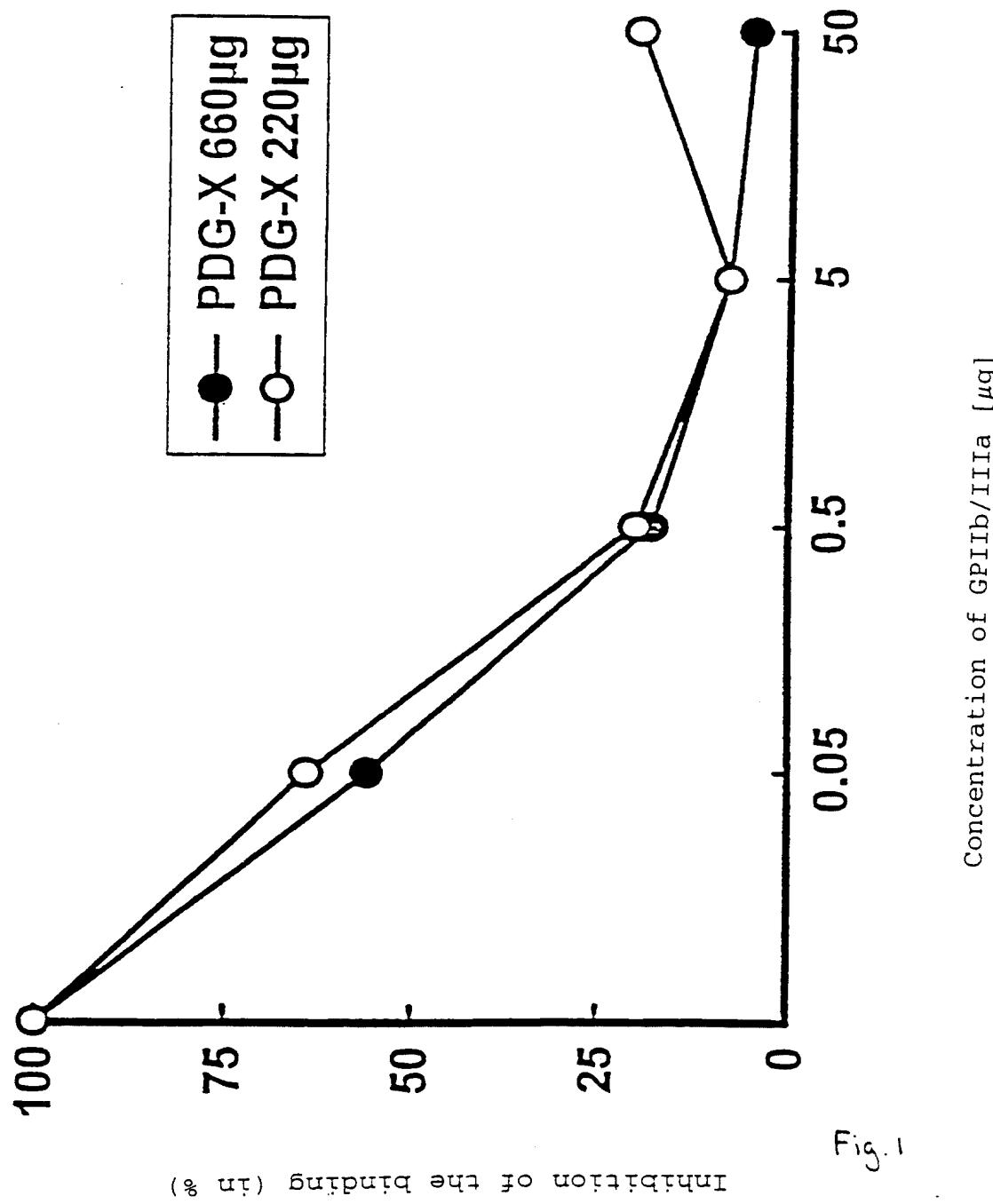
10. Nucleic acid from Claim 9, which furthermore encompasses a CDR1 and/or CDR2 region selected from a nucleotide sequence which encodes the amino acid sequences shown in Tab. 7a or b or an amino acid sequence which is at least 80% homologous thereto.
11. Vector, characterized in that it
 - (a) contains at least one copy of a nucleic acid according to one of Claims 1 to 3 and/or at least one copy of a nucleic acid according to one of Claims 4 to 6 or
 - (b) contains at least one copy of a nucleic acid according to Claim 7 or 8 and/or at least one copy of a nucleic acid according to Claim 9 or 10.
12. Cell, characterized in that it
 - (a) expresses a nucleic acid according to one of Claims 1 to 3 and/or a nucleic acid according to one of Claims 4 to 6 or
 - (b) a nucleic acid according to Claim 7 or 8 and/or a nucleic acid according to Claim 9 or 10.
13. Polypeptide, characterized in that it
 - (a) is encoded by a nucleic acid according to one of Claims 1 to 3 and/or a nucleic acid according to one of Claims 4 to 8 or
 - (b) by a nucleic acid according to Claim 7 or 8 and/or a nucleic acid according to Claim 9 or 10.

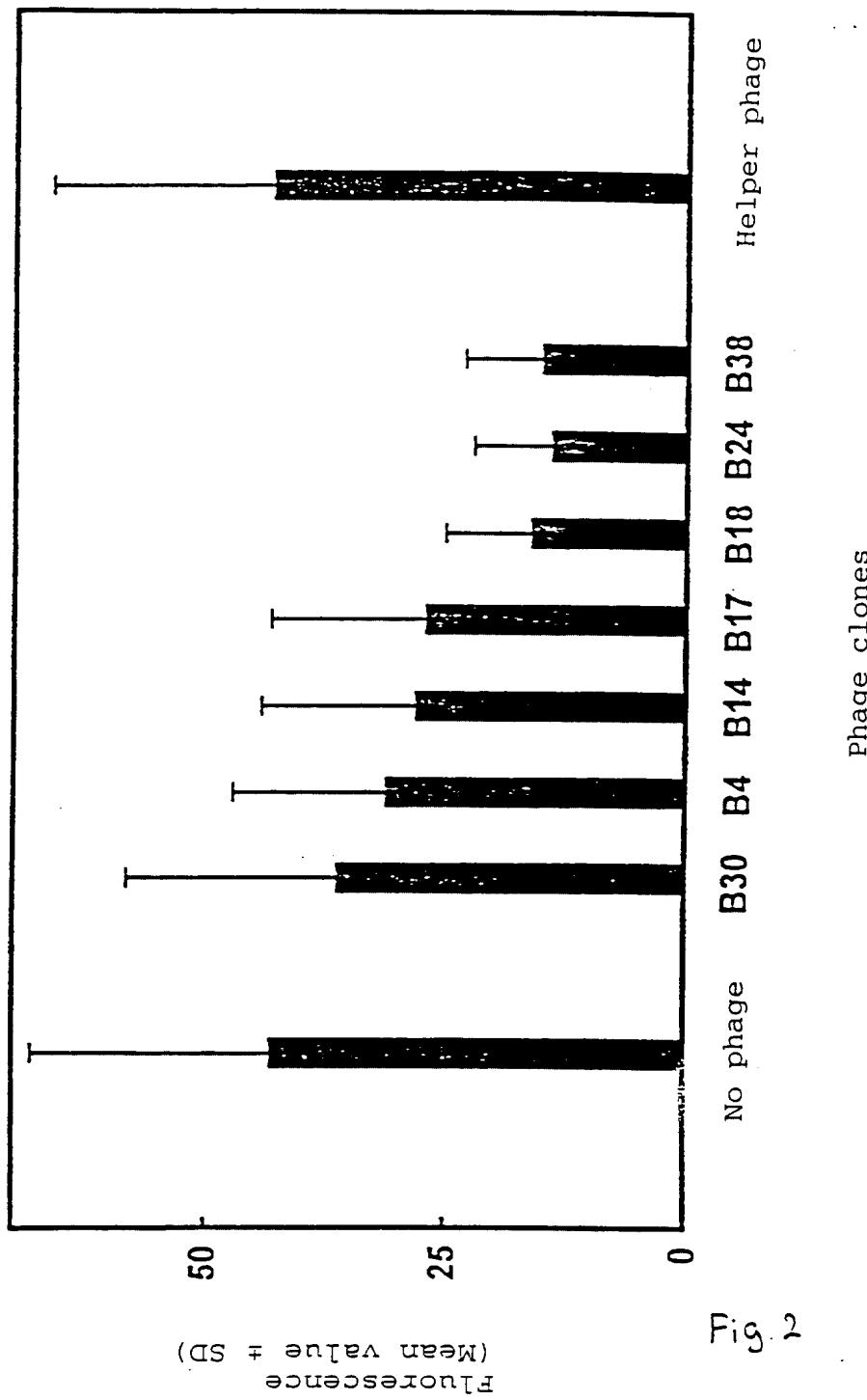
14. Polypeptide according to Claim 13, characterized in that it comprises the variable domain of the H chain and/or the variable domain of the L chain of a human antibody.
15. Polypeptide according to Claim 14, characterized in that it comprises both the variable domain of the H chain and the variable domain of the L chain.
16. Polypeptide according to one of Claims 13 to 15, characterized in that it is coupled to a labelling group or a toxin.
17. Antibody against a polypeptide according to one of Claims 13 to 16.
18. Antibody according to Claim 17, characterized in that it is directed against the CDR3 region of the heavy and/or light antibody chain of the polypeptide.
19. Pharmaceutical composition which comprises, as the active component, a nucleic acid according to one of Claims 1 to 10, a vector according to Claim 11, a cell according to Claim 12, a polypeptide according to one of Claims 13 to 16 or an antibody according to either Claim 17 or 18, where appropriate together with other active components and pharmaceutically customary adjuvants, additives or excipients.

20. Use of a nucleic acid according to one of Claims 1 to 10, of a vector according to Claim 11, of a cell according to Claim 12, of a polypeptide according to one of Claims 13 to 16, of an antibody according to Claim 17 or 18, or of a pharmaceutical composition according to Claim 19 for preparing an agent for the diagnosis or for the treatment or prevention of AITP.

21. Use of a nucleic acid according to one of Claims 1 to 10, of a vector according to Claim 11, of a cell according to Claim 12, of a polypeptide according to one of Claims 13 to 16, or of a pharmaceutical composition according to Claim 19 for preparing an agent for exerting an effect on the binding of fibrinogen to blood platelets.
22. Use according to Claim 21 for preparing an agent for modulating blood coagulation, in particular for dissolving thrombi and/or for preventing the formation of thrombi.
23. Process for isolating phagemid clones which express nucleic acids which encode autoantibodies against GPIIb/IIIa or encode antiidiotypic antibodies which are directed against these autoantibodies, characterized in that a phagemid library is prepared from lymphocytes obtained from a healthy human donor and the desired phagemid clones are isolated by affinity selection comprising negative and positive selection steps.
24. Process according to Claim 23, characterized in that antibody-encoding nucleic acids are isolated from the clones.
25. Process according to Claim 23 or 24, characterized in that the antibody-encoding nucleic acids are used for expressing recombinant antibody chains, or derivatives or fragments thereof.

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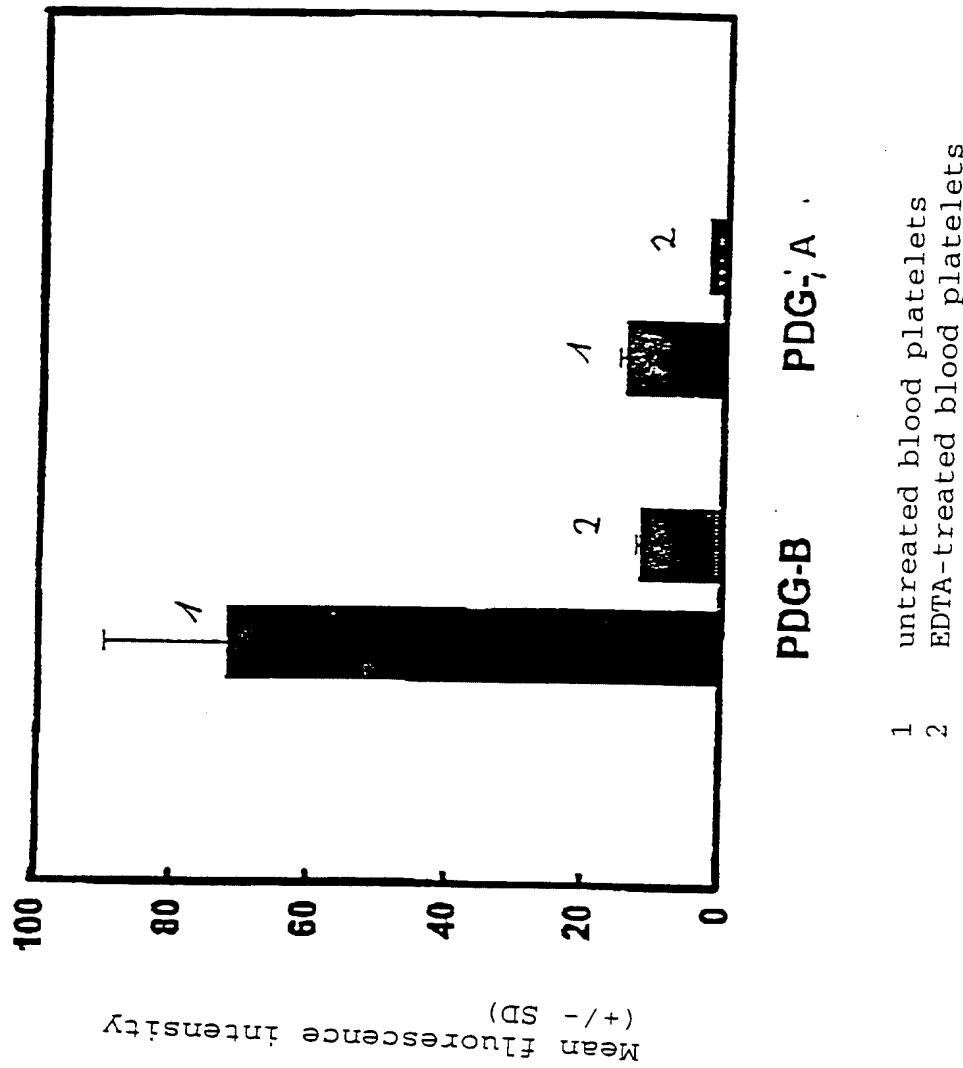
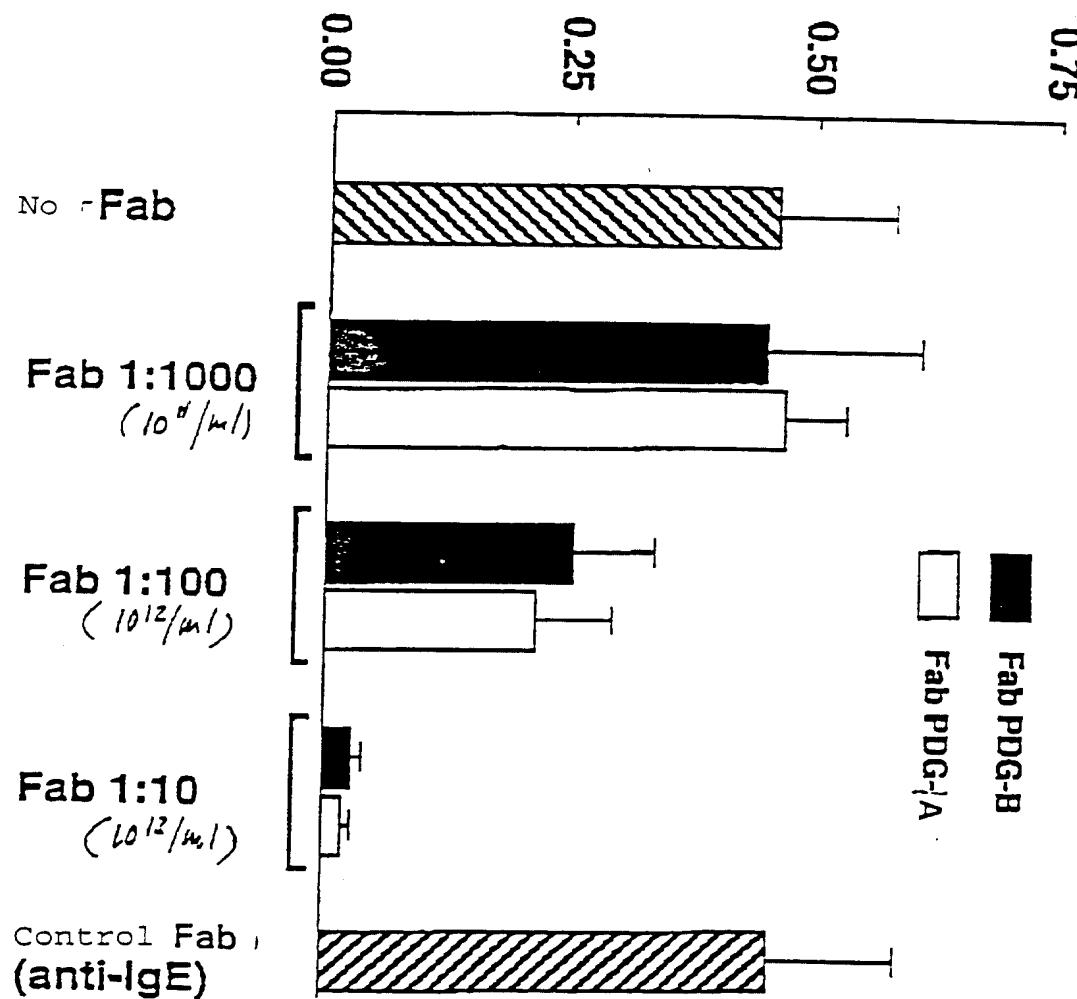


Fig. 3

Fibrinogen binding
(mean O/D +/- SD)

Fig. 4



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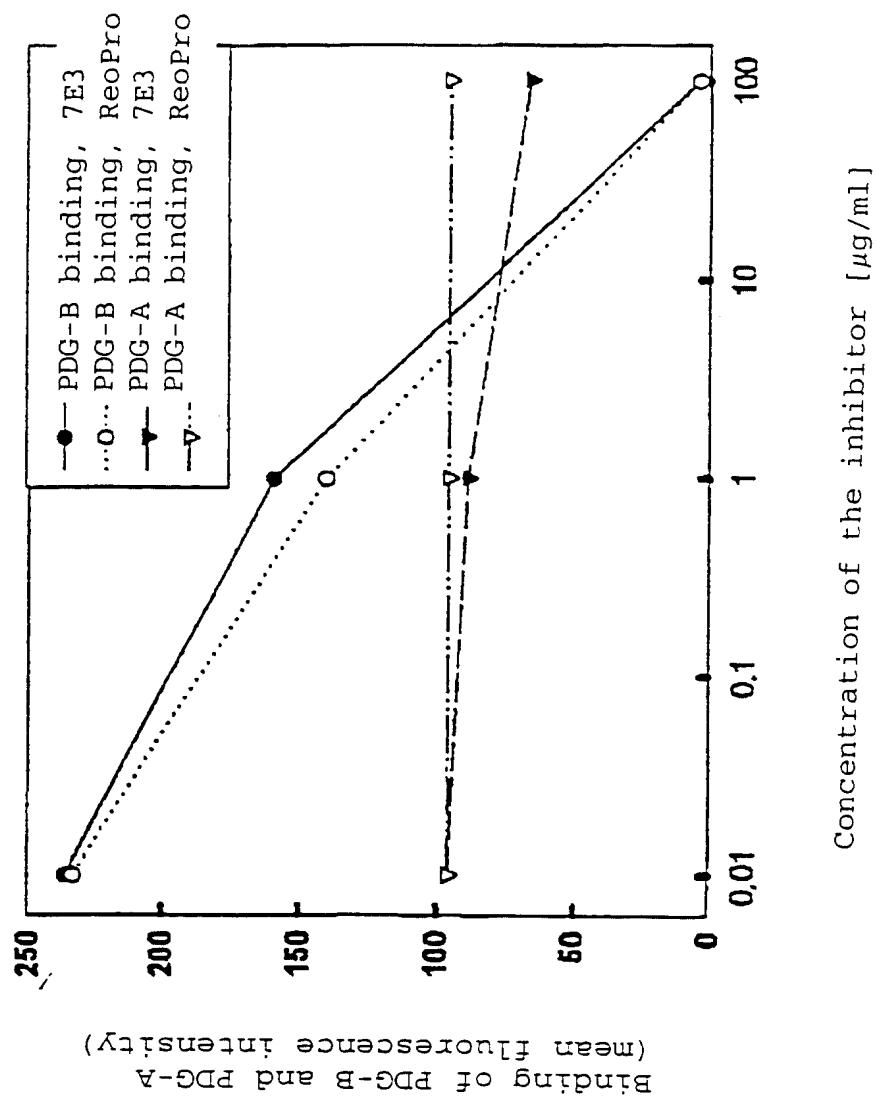


Fig. 5

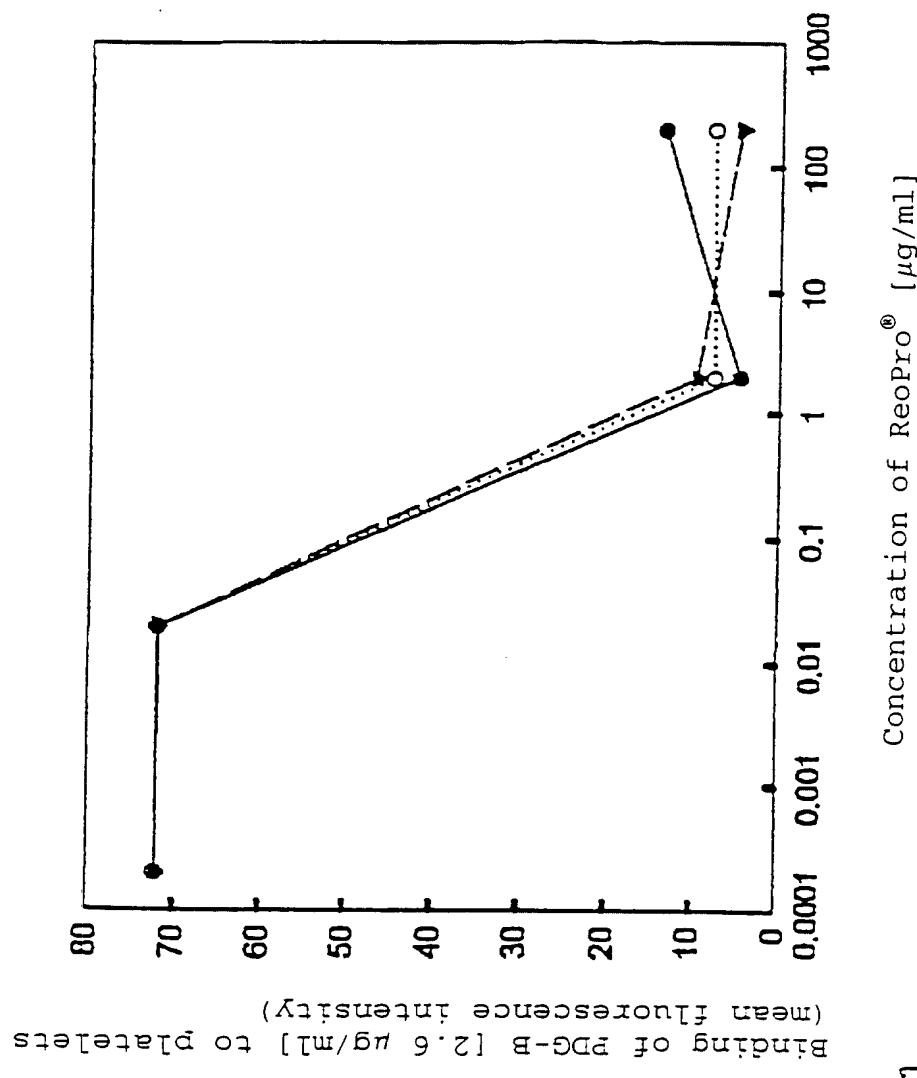


Fig. 6

—●— 5 min incubation with PDG-B, then addition of ReoPro[®] for 1.5 h
····○··· Incubation with ReoPro[®] for 5 min, then addition of PDG for 1.5 h
—▽— Incubation with PDG-B and ReoPro, added simultaneously

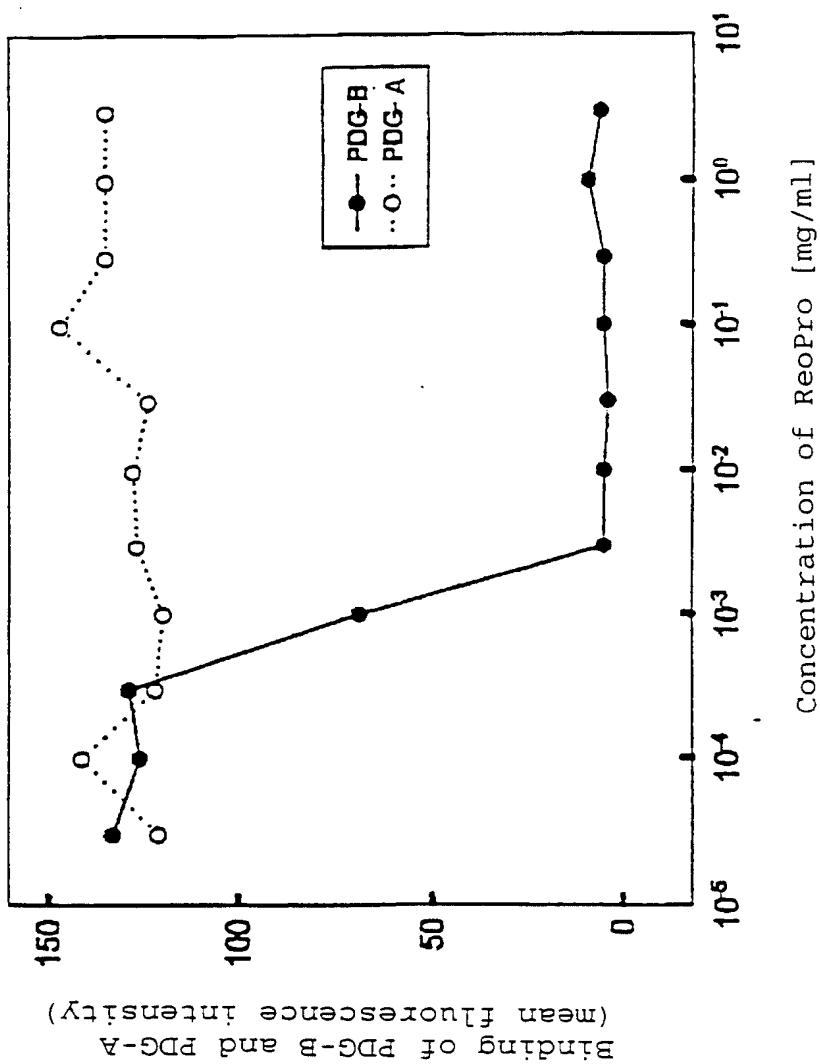


Fig. 7